



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C12N 15/31, C07K 14/315, A61K 39/09, C12N 1/21	A2	(11) International Publication Number: WO 99/42588 (43) International Publication Date: 26 August 1999 (26.08.99)
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(54) Title: GROUP B STREPTOCOCCUS ANTIGENS		
(57) Abstract Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.		

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GROUP B STREPTOCOCCUS ANTIGENS

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FIELD OF THE INVENTION

The present invention is related to antigens, more particularly protein antigens of group B streptococcus (GBS) bacterial pathogen which are useful as vaccine components for therapy and/or prophylaxis.

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BACKGROUND OF THE INVENTION

Streptococcus are gram (+) bacteria that are differentiated by group specific carbohydrate antigens A through O found on their cell surface. Streptococcus groups are further distinguished by type-specific capsular polysaccharide antigens. Several serotypes have been identified for the Group B streptococcus (GBS) : Ia, Ib, II, III, IV, V, VI, VII and VIII. GBS also contains antigenic proteins known as "C-proteins" (alpha, beta, gamma and delta), some of which have been cloned.

25

Although GBS is a common component of the normal human vaginal and colonic flora this pathogen has long been recognized as a major cause of neonatal sepsis and meningitis, late-onset meningitis in infants, postpartum endometritis as well as mastitis in dairy herds. Expectant mothers exposed to GBS are at risk of postpartum infection and may transfer the infection to their baby as the child passes through the birth canal. Although the organism is sensitive to antibiotics, the high attack rate and rapid onset of sepsis in neonates and meningitis in infants results in high morbidity and mortality.

To find a vaccine that will protect individuals from GBS infection, researches have turned to the type-specific antigens. Unfortunately these polysaccharides have proven to
5 be poorly immunogenic in humans and are restricted to the particular serotype from which the polysaccharide originates. Further, capsular polysaccharide elicit a T cell independent response i.e. no IgG production. Consequently capsular polysaccharide antigens are unsuitable
10 as a vaccine component for protection against GBS infection.

Others have focused on the C-protein beta antigen which demonstrated immunogenic properties in mice and rabbit models. This protein was found to be unsuitable as a human
15 vaccine because of its undesirable property of interacting with high affinity and in a non-immunogenic manner with the Fc region of human IgA. The C-protein alpha antigen is rare in type III serotypes of GBS which is the serotype responsible for most GBS mediated conditions and is
20 therefore of little use as a vaccine component.

Therefore there remains an unmet need for GBS antigens that may be used as vaccine components for the prophylaxis and/or
25 therapy of GBS infection.

SUMMARY OF THE INVENTION

30 According to one aspect, the present invention provides an isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide comprising a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
35 SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
5 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments,
analogs or derivatives thereof.

In other aspects, there is provided vectors comprising
polynucleotides of the invention operably linked to an
10 expression control region, as well as host cells transfected
with said vectors and methods of producing polypeptides
comprising culturing said host cells under conditions
suitable for expression.

15 In yet another aspect, there is provided novel polypeptides
encoded by polynucleotides of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

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Figure 1a is the DNA sequence of clone 1 (SEQ ID NO :1) with
corresponding amino acid sequences for open reading frames;
figure 1b is the amino acid sequence SEQ ID NO: 2;
figure 1c is the amino acid sequence SEQ ID NO: 3;
25 figure 1d is the amino acid sequence SEQ ID NO: 4;
figure 1e is the amino acid sequence SEQ ID NO: 5;
figure 1f is the amino acid sequence SEQ ID NO: 6;

Figure 2a is the DNA sequence of clone 2 (SEQ ID NO :7) with
30 corresponding amino acid sequences for open reading frames;
figure 2b is the amino acid sequence SEQ ID NO: 8;
figure 2c is the amino acid sequence SEQ ID NO: 9;
figure 2d is the amino acid sequence SEQ ID NO:10;
figure 2e is the amino acid sequence SEQ ID NO:11;
35 figure 2f is the amino acid sequence SEQ ID NO:12;

Figure 3a is the DNA sequence of clone 3 (SEQ ID NO :13)
with corresponding amino acid sequences for open reading
frames;

figure 3b is the amino acid sequence SEQ ID NO:14;

5 figure 3c is the amino acid sequence SEQ ID NO:15;

figure 3d is the amino acid sequence SEQ ID NO:16;

figure 3e is the amino acid sequence SEQ ID NO:17;

figure 3f is the amino acid sequence SEQ ID NO:18;

figure 3g is the amino acid sequence SEQ ID NO:19;

10 figure 3h is the amino acid sequence SEQ ID NO:20;

figure 3i is the amino acid sequence SEQ ID NO:21;

Figure 4a is the DNA sequence of clone 4 (SEQ ID NO :22)
with corresponding amino acid sequences for open reading
15 frames;

figure 4b is the amino acid sequence SEQ ID NO:23;

figure 4c is the amino acid sequence SEQ ID NO:24;

figure 4d is the amino acid sequence SEQ ID NO:25;

20 figure 4e is the amino acid sequence SEQ ID NO:26;

Figure 5a is the DNA sequence of clone 5 (SEQ ID NO :27)
with corresponding amino acid sequences for open reading
frames;

figure 5b is the amino acid sequence SEQ ID NO:28;

25 figure 5c is the amino acid sequence SEQ ID NO:29;

figure 5d is the amino acid sequence SEQ ID NO:30;

figure 5e is the amino acid sequence SEQ ID NO:31;

Figure 6a is the DNA sequence of clone 6 (SEQ ID NO :32) ;

30 figure 6b is the amino acid sequence SEQ ID NO:33;

figure 6c is the amino acid sequence SEQ ID NO:34;

figure 6d is the amino acid sequence SEQ ID NO:35;

figure 6e is the amino acid sequence SEQ ID NO:36;

35 Figure 7a is the DNA sequence of clone 7 (SEQ ID NO :37);

figure 7b is the amino acid sequence SEQ ID NO:38;

figure 7c is the amino acid sequence SEQ ID NO:39;

figure 7d is the amino acid sequence SEQ ID NO:40;

figure 7e is the amino acid sequence SEQ ID NO:41;

- 5 Figure 8 is the DNA sequence of a part of clone 7 including a signal sequence (SEQ ID NO :42);

Figure 9 is the DNA sequence of a part of clone 7 without a signal sequence (SEQ ID NO :43);

- 10 Figure 9a is the amino acid sequence (SEQ ID NO:44);

Figure 10 represents the distribution of anti-GBS ELISA titers in sera from CD-1 mice immunized with recombinant GBS protein corresponding to the SEQ ID NO:39.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to novel antigenic polypeptides of group B streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments,
analogues or derivatives thereof.

A preferred embodiment of the invention includes SEQ ID NO :39 and SEQ ID NO:44.

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A further preferred embodiment of the invention is SEQ ID NO :39.

A further preferred embodiment of the invention is SEQ ID NO :44.

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As used herein, "fragments", "derivatives" or "analogues" of the polypeptides of the invention include those polypeptides in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably conserved) and which may be natural or unnatural.

30

The terms «fragments», «derivatives» or «analogues» of polypeptides of the present invention also include polypeptides which are modified by addition, deletion,

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substitution of amino acids provided that the polypeptides retain the capacity to induce an immune response.

- By the term «conserved amino acid» is meant a substitution of one or more amino acids for another in which the antigenic determinant (including its secondary structure and hydropathic nature) of a given antigen is completely or partially conserved in spite of the substitution.
- For example, one or more amino acid residues within the sequence can be substituted by another amino acid of a similar polarity, which acts as a functional equivalent, resulting in a silent alteration. Substitutes for an amino acid within the sequence may be selected from other members of the class to which the amino acid belongs. For example, the nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan and methionine. The polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine and glutamine. The positively charged (basic) amino acids include arginine, lysine and histidine. The negatively charged (acidic) amino acids include aspartic acid and glutamic acid.
- Preferably, derivatives and analogs of polypeptides of the invention will have about 70% identity with those sequences illustrated in the figures or fragments thereof. That is, 70% of the residues are the same. More preferably polypeptides will have greater than 95% homology. In another preferred embodiment, derivatives and analogs of polypeptides of the invention will have fewer than about 20 amino acid residue substitutions, modifications or deletions and more preferably less than 10. Preferred substitutions are those known in the art as conserved i.e. the substituted residues share physical or chemical properties such as hydrophobicity, size, charge or functional groups.

Furthermore, in those situations where amino acid regions are found to be polymorphic, it may be desirable to vary one or more particular amino acids to more effectively mimic the different epitopes of the different GBS strains.

Also included are polypeptides which have fused thereto other compounds which alter the polypeptides biological or pharmacological properties i.e. polyethylene glycol (PEG) to increase half-life; leader or secretory amino acid sequences for ease of purification; prepro- and pro- sequences; and (poly)saccharides.

Moreover, the polypeptides of the present invention can be modified by terminal -NH₂ acylation (eg. by acetylation, or thioglycolic acid amidation, terminal carboxy amidation, e.g. with ammonia or methylamine) to provide stability, increased hydrophobicity for linking or binding to a support or other molecule.

Also contemplated are hetero and homo polypeptide multimers of the polypeptide fragments, analogues and derivatives. These polymeric forms include, for example, one or more polypeptides that have been cross-linked with cross-linkers such as avidin/biotin, gluteraldehyde or dimethylsuperimide. Such polymeric forms also include polypeptides containing two or more tandem or inverted contiguous sequences, produced from multicistronic mRNAs generated by recombinant DNA technology.

Preferably, a fragment, analog or derivative of a polypeptide of the invention will comprise at least one antigenic region i.e. at least one epitope.

In order to achieve the formation of antigenic polymers (i.e. synthetic multimers), polypeptides may be utilized having bishaloacetyl groups, nitroarylhalides, or the like,

where the reagents being specific for thio groups.
Therefore, the link between two mercapto groups of the
different peptides may be a single bond or may be composed
of a linking group of at least two, typically at least four,
5 and not more than 16, but usually not more than about 14
carbon atoms.

In a particular embodiment, polypeptide fragments, analogs
and derivatives of the invention do not contain a methionine
10 (Met) starting residue. Preferably, polypeptides will not
incorporate a leader or secretory sequence (signal
sequence). The signal portion of a polypeptide of the
invention may be determined according to established
molecular biological techniques. In general, the
15 polypeptide of interest may be isolated from a GBS culture
and subsequently sequenced to determine the initial residue
of the mature protein and therefor the sequence of the
mature polypeptide.

20 According to another aspect, there is provided vaccine
compositions comprising one or more GBS polypeptides of the
invention in admixture with a pharmaceutically acceptable
carrier diluent or adjuvant.

25 Suitable adjuvants include oils i.e. Freund's complete or
incomplete adjuvant; salts i.e. $\text{AlK}(\text{SO}_4)_2$, $\text{AlNa}(\text{SO}_4)_2$,
 $\text{AlNH}_4(\text{SO}_4)_2$, $\text{Al}(\text{OH})_3$, AlPO_4 , silica, kaolin; saponin
derivative; carbon polynucleotides i.e. poly IC and poly AU
and also detoxified cholera toxin (CTB) and E.coli heat
30 labile toxin for induction of mucosal immunity. Preferred
adjuvants include QuilATM, AlhydrogelTM and AdjuphosTM.
Vaccines of the invention may be administered parenterally
by injection, rapid infusion, nasopharyngeal absorption,
dermoabsorption, or bucal or oral.

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- Vaccine compositions of the invention are used for the treatment or prophylaxis of *streptococcus* infection and/or diseases and symptoms mediated by *streptococcus* infection,
- 5 in particular group A *streptococcus* (*pyogenes*), group B *streptococcus* (GBS or *agalactiae*), *dysgalactiae*, *uberis*, *nocardia* as well as *Staphylococcus aureus*. General information about *Streptococcus* is available in Manual of Clinical Microbiology by P.R.Murray et al. (1995, 6th Edition,
- 10 ASM Press, Washington, D.C.). More particularly group B *streptococcus*, *agalactiae*. In a particular embodiment vaccines are administered to those individuals at risk of GBS infection such as pregnant women and infants for sepsis, meningitis and pneumonia as well as immunocompromised
- 15 individuals such as those with diabetes, liver disease or cancer. Vaccines may also have veterinary applications such as for the treatment of mastitis in cattle which is mediated by the above mentioned bacteria as well as *E.coli*.
- 20 The vaccine of the present invention can also be used for the manufacture of a medicament used for the treatment or prophylaxis of *streptococcus* infection and/or diseases and symptoms mediated by *streptococcus* infection, in particular group A *streptococcus* (*pyogenes*), group B *streptococcus* (GBS
- 25 or *agalactiae*), *dysgalactiae*, *uberis*, *nocardia* as well as *Staphylococcus aureus*. More particularly group B *streptococcus*, *agalactiae*.

- Vaccine compositions are preferably in unit dosage form of
- 30 about 0.001 to 100 µg/kg (antigen/body weight) and more preferably 0.01 to 10 µg/kg and most preferably 0.1 to 1 µg/kg 1 to 3 times with an interval of about 1 to 12 weeks intervals between immunizations, and more preferably 1 to 6

weeks.

According to another aspect, there is provided polynucleotides encoding polypeptides of group B streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO : 42) and 9 (SEQ ID NO : 43) which correspond to the open reading frames, encoding polypeptides of the invention.

Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO : 42) and 9 (SEQ ID NO : 43) and fragments, analogues and derivatives thereof.

More preferred polynucleotides of the invention are those illustrated in Figures 7 (SEQ ID NO : 37), 8 (SEQ ID NO : 42) and 9 (SEQ ID NO : 43).

Most preferred polynucleotides of the invention are those illustrated in Figures 8 (SEQ ID NO : 42) and 9 (SEQ ID NO :

43).

It will be appreciated that the polynucleotide sequences illustrated in the figures may be altered with degenerate
5 codons yet still encode the polypeptides of the invention.

Due to the degeneracy of nucleotide coding sequences, other polynucleotide sequences which encode for substantially the same polypeptides of the present invention may be used in
10 the practice of the present invention. These include but are not limited to nucleotide sequences which are altered by the substitution of different codons that encode the same amino acid residue within the sequence, thus producing a silent change.

15 Accordingly the present invention further provides polynucleotides which hybridize to the polynucleotide sequences herein above described (or the complement sequences thereof) having 50% and preferably at least 70%
20 identity between sequences. More preferably polynucleotides are hybridizable under stringent conditions i.e. having at least 95% identity and most preferably more than 97% identity.

25 By capable of hybridizing under stringent conditions is meant annealing of a nucleic acid molecule to at least a region of a second nucleic acid sequence (whether as cDNA, mRNA, or genomic DNA) or to its complementary strand under standard conditions, e.g. high temperature and/or low salt
30 content, which tend to disfavor hybridization of noncomplementary nucleotide sequences. A suitable protocol is described in Maniatis T. et al., Molecular cloning : A Laboratory Manual, Cold Springs Harbor Laboratory, 1982, which is herein incorporated by reference.

35 In a further aspect, polynucleotides encoding polypeptides

of the invention, or fragments, analogs or derivatives thereof, may be used in a DNA immunization method. That is, they can be incorporated into a vector which is replicable and expressible upon injection thereby producing the antigenic polypeptide in vivo. For example polynucleotides may be incorporated into a plasmid vector under the control of the CMV promoter which is functional in eukaryotic cells. Preferably the vector is injected intramuscularly.

10

According to another aspect, there is provided a process for producing polypeptides of the invention by recombinant techniques by expressing a polynucleotide encoding said polypeptide in a host cell and recovering the expressed polypeptide product. Alternatively, the polypeptides can be produced according to established synthetic chemical techniques i.e. solution phase or solid phase synthesis of oligopeptides which are ligated to produce the full polypeptide (block ligation).

20

For recombinant production, host cells are transfected with vectors which encode the polypeptide, and then cultured in a nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes. Suitable vectors are those that are viable and replicable in the chosen host and include chromosomal, non-chromosomal and synthetic DNA sequences e.g. bacterial plasmids, phage DNA, baculovirus, yeast plasmids, vectors derived from combinations of plasmids and phage DNA. The polypeptide sequence may be incorporated in the vector at the appropriate site using restriction enzymes such that it is operably linked to an expression control region comprising a promoter, ribosome binding site (consensus region or Shine-Dalgarno sequence), and optionally an operator (control element). One can select individual components of the expression control region that are appropriate for a given

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host and vector according to established molecular biology principles (Sambrook et al, Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor, N.Y., 1989 incorporated herein by reference). Suitable promoters include but are not
5 limited to LTR or SV40 promoter, *E.coli* lac, tac or trp promoters and the phage lambda P_L promoter. Vectors will preferably incorporate an origin of replication as well as selection markers i.e. ampicillin resistance gene. Suitable bacterial vectors include pET, pQE70, pQE60, pQE-9, pbs, pD10 phagescript, psiX174, pbluescript SK, pbsks, pNH8A,
10 pNH16a, pNH18A, pNH46A, ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 and eukaryotic vectors pBlueBacIII, pWLNEO, pSV2CAT, pOG44, pXT1, pSG, pSVK3, pBPV, pMSG and pSVL. Host cells may be bacterial i.e. *E.coli*, *Bacillus subtilis*,
15 *Streptomyces*; fungal i.e. *Aspergillus niger*, *Aspergillus nidulins*; yeast i.e. *Saccharomyces* or eukaryotic i.e. CHO, COS.

Upon expression of the polypeptide in culture, cells are
20 typically harvested by centrifugation then disrupted by physical or chemical means (if the expressed polypeptide is not secreted into the media) and the resulting crude extract retained to isolate the polypeptide of interest. Purification of the polypeptide from culture media or lysate
25 may be achieved by established techniques depending on the properties of the polypeptide i.e. using ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, hydroxylapatite
30 chromatography and lectin chromatography. Final purification may be achieved using HPLC.

The polypeptide may be expressed with or without a leader or secretion sequence. In the former case the leader may be
35 removed using post-translational processing (see US

4,431,739; 4,425,437; and 4,338,397 incorporated herein by reference) or be chemically removed subsequent to purifying the expressed polypeptide.

- 5 According to a further aspect, the GBS polypeptides of the invention may be used in a diagnostic test for streptococcus infection in particular GBS infection. Several diagnostic methods are possible, for example detecting streptococcus organism in a biological sample, the following procedure may
- 10 be followed:
- a) obtaining a biological sample from a patient;
 - b) incubating an antibody or fragment thereof reactive with a GBS polypeptide of the invention with the biological sample to form a mixture; and
 - 15 c) detecting specifically bound antibody or bound fragment in the mixture which indicates the presence of streptococcus.

- Alternatively, a method for the detection of antibody
- 20 specific to a streptococcus antigen in a biological sample containing or suspected of containing said antibody may be performed as follows:
- a) isolating a biological sample from a patient;
 - b) incubating one or more GBS polypeptides of the
 - 25 invention or fragments thereof with the biological sample to form a mixture; and
 - c) detecting specifically bound antigen or bound fragment in the mixture which indicates the presence of antibody specific to streptococcus.

- 30 One of skill in the art will recognize that this diagnostic test may take several forms, including an immunological test such as an enzyme-linked immunosorbent assay (ELISA), a radioimmunoassay or a latex agglutination assay, essentially
- 35 to determine whether antibodies specific for the protein are present in an organism.

The DNA sequences encoding polypeptides of the invention may also be used to design DNA probes for use in detecting the presence of streptococcus in a biological sample suspected of containing such bacteria. The detection method of this invention comprises:

- a) isolating the biological sample from a patient;
- b) incubating one or more DNA probes having a DNA sequence encoding a polypeptide of the invention or fragments thereof with the biological sample to form a mixture; and
- c) detecting specifically bound DNA probe in the mixture which indicates the presence of streptococcus bacteria.

The DNA probes of this invention may also be used for detecting circulating streptococcus i.e. GBS nucleic acids in a sample, for example using a polymerase chain reaction, as a method of diagnosing streptococcus infections. The probe may be synthesized using conventional techniques and may be immobilized on a solid phase, or may be labeled with a detectable label. A preferred DNA probe for this application is an oligomer having a sequence complementary to at least about 6 contiguous nucleotides of the GBS polypeptides of the invention.

Another diagnostic method for the detection of streptococcus in a patient comprises:

- a) labeling an antibody reactive with a polypeptide of the invention or fragment thereof with a detectable label;
- b) administering the labeled antibody or labeled fragment to the patient; and
- c) detecting specifically bound labeled antibody or labeled fragment in the patient which indicates the presence of streptococcus.

A further aspect of the invention is the use of the GBS

polypeptides of the invention as immunogens for the production of specific antibodies for the diagnosis and in particular the treatment of streptococcus infection. Suitable antibodies may be determined using appropriate screening methods, for example by measuring the ability of a particular antibody to passively protect against streptococcus infection in a test model. One example of an animal model is the mouse model described in the examples herein. The antibody may be a whole antibody or an antigen-binding fragment thereof and may in general belong to any immunoglobulin class. The antibody or fragment may be of animal origin, specifically of mammalian origin and more specifically of murine, rat or human origin. It may be a natural antibody or a fragment thereof, or if desired, a recombinant antibody or antibody fragment. The term recombinant antibody or antibody fragment means antibody or antibody fragment which were produced using molecular biology techniques. The antibody or antibody fragments may be polyclonal, or preferably monoclonal. It may be specific for a number of epitopes associated with the GBS polypeptides but is preferably specific for one.

EXAMPLE 1 Murine model of lethal Group B Streptococcus (GBS) infection

The mouse model of GBS infection is described in detail in Lancefield et al (J Exp Med 142:165-179,1975). GBS strain C388/90 (Clinical isolate obtained in 1990 from the cephalorachidian fluid of a patient suffering from meningitis, Children's Hospital of Eastern Ontario, Ottawa, Canada) and NCS246 (National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Edmonton, Canada) were respectively serotyped as type Ia/c and type II/R.

To increase their virulence, the GBS strains C388/90 (serotype Ia/c) and NCS 246 (serotype II/R) were serially passaged through mice as described previously (Lancefield et al. J Exp Med 142:165-179, 1975). Briefly, the increase of virulence was monitored using intraperitoneal inoculations of serial dilutions of a subculture in Todd-Hewitt broth obtained from either the blood or spleen of infected mice. After the last passage, infected blood samples were used to inoculate Todd-Hewitt broth. After an incubation of 2 hours at 37°C with 7% CO₂, glycerol at a final concentration of 10% (v/v) was added to the culture. The culture was then aliquoted and stored at -80° C for use in GBS challenge experiments. The number of cfu of GBS present in these frozen samples was determined. The bacterial concentration necessary to kill 100% (LD100) of the 18 weeks old mice were determined to be 3.5X10⁵ and 1.1X10⁵ respectively for GBS strain C388/90 and NCS246, which corresponded to a significant increase in virulence for both strains. Indeed, the LD100 recorded before the passages for these two strains was higher than 10⁹ cfu.

In a bacterial challenge, a freshly thawed aliquot of a virulent GBS strain was adjusted to the appropriate bacterial concentration using Todd-Hewitt broth and 1ml was injected intraperitoneally to each female CD-1 mouse. The mice used for the passive protection experiments were 6 to 8 weeks old, while the ones used for the active protection experiments were approximately 18 weeks old at the time of the challenge. All inocula were verified by colony counts. Animals were observed for any sign of infection four times daily for the first 48 h after challenge and then daily for the next 12 days. At the end of that period, blood samples were obtained from the survivors and frozen at -20°C. The spleen obtained from each mouse that survived the challenge was cultured in order to identify any remaining GBS.

EXAMPLE 2 Immunization and protection in mice with formaldehyde killed whole GBS cells

- 5 Formaldehyde killed GBS whole cells were prepared according to the procedures described in Lancefield et al (J Exp Med 142:165-179,1975). Briefly, an overnight culture on sheep blood agar plates (Quelab Laboratories, Montreal, Canada) of a GBS strain was washed twice in PBS buffer (phosphate buffered-saline, pH7.2), adjusted to approximately 3×10^9 cfu/mL and incubated overnight in PBS containing 0.3% (v/v) formaldehyde. The killed GBS suspension was washed with PBS and kept frozen at -80°C .
- 15 Female CD-1 mice, 6 to 8 weeks old (Charles River, St-Constant, Québec, Canada), were injected subcutaneously three times at two weeks interval with 0.1 ml of formaldehyde killed cells of GBS strain C388/90 ($\sim 6 \times 10^7$ GBS), or 0.1 ml of PBS for the control group. On the day before the immunization, AlhydrogelTM (Superfos Biosector, Frederikssund, Denmark) at a final concentration of 0.14 mg or 0.21 mg of Al, was added to these preparations and incubated overnight at 4°C with agitation. Serum samples were obtained from each mouse before the beginning of the immunization protocol and two weeks after the last injection. The sera were frozen at -20°C .

- Eight mice in each control group injected with PBS and the group immunized with formaldehyde killed whole cells GBS strain C388/90 (Ia/c) were challenged with 1.5×10^4 cfu of GBS strain C388/90 (Ia/c) one week after the third injection. All mice immunized with the formaldehyde killed GBS whole cells survived the homologous challenge while, within 5 days after the challenge, only 4 out of the 8 mice injected with PBS survived from the infection. In order to increase the mortality rate in the control groups, the

bacterial suspension had to be adjusted according to the age of the mice at the time of the bacterial challenge. In subsequent challenge experiments, when mice were older than 15 weeks, the bacterial inoculum was increased to

5 concentrations between 3.0×10^5 and 2.5×10^6 cfu.

Table 1 Immunization of CD1 mice with formaldehyde killed whole cells of GBS and subsequent homologous challenge [strain C388/90 (Ia/c)] and heterologous challenge [strain NCS246 (II/R)].

antigenic preparations used for immunization ¹	number of living mice 14 days after the bacterial challenge (% Survival)	
	homologous challenge: strain C388/90 (Ia/c)	heterologous challenge: strain NCS246 (II/R)
1st infection		
formaldehyde killed cells of GBS strain C388/90 (Ia/c) ²	8/8 (100) ³	n.d. ⁵
control PBS	4/8 (50)	n.d.
2nd infection		
formaldehyde killed cells of GBS strain C388/90 (Ia/c)	6/6 (100) ⁴	0/6 (0) ⁶
control PBS	2/6 (33)	0/6 (0)

¹ alhydrogel™ at a final concentration of 0.14 mg or 0.21mg of AI was used;

² approximately 6×10^7 cfu;

³ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 1.5×10^4 cfu;

⁴ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2.1×10^5 cfu;

⁵ not done;

⁶ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS NCS246 (II/R) suspension adjusted to 1.2×10^5 cfu.

In another experiment, one group of 12 mice corresponding to a control group was injected with PBS, while a second group of 12 mice was immunized with formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Six mice from each of these two groups were challenged with 2.1×10^6 cfu of the GBS strain C388/90 (Ia/c) (Table I). As the first challenge experiment, all mice immunized with the GBS strain C388/90 (Ia/c) survived the homologous challenge. Only two out of the 6 mice injected with PBS survived the infection.

The remaining 6 mice in both groups were then used one week later to verify whether this antigenic preparation could confer cross protection against strain NCS246 (II/R) which produce a serologically distinct capsule. None of the mice infected with this second GBS strain survived the infection. The later result suggested that most of the protective immune response induced by formaldehyde killed strain C388/90 is directed against the capsular polysaccharide and that it could be restricted to strains of that particular serotype. These results clearly indicated that this particular model of infection can be efficiently used to study the protection conferred by vaccination.

15 EXAMPLE 3 Immunization of rabbit with formaldehyde killed whole GBS cells and passive protection in mice

A New Zealand rabbit (2.5 kg, Charles River, St-Constant, Québec, Canada) was immunized with formaldehyde killed cells of GBS strain C388/90 (Ia/c) to obtain hyperimmune serum. This rabbit was injected subcutaneously three times at three weeks interval with approximately 1.5×10^9 cfu of formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Freund's complete adjuvant (Gibco BRL Life Technologies, Grand Island, New York) was used as the adjuvant for the first immunization, while Freund's incomplete adjuvant (Gibco BRL) was used for the following two injections. Serum samples were obtained before the beginning of the immunization protocol and two weeks after the last injection. The sera were frozen at -20°C .

The ability of this particular rabbit hyperimmune serum to passively protect mice against a lethal infection with GBS

was also evaluated. Intraperitoneal injection of mice with either 15 or 25 μ L of hyperimmune rabbit serum 18 hours before the challenge protected 4 out of 5 mice (80%) against the infection. Comparatively, survival rates lower than 20% were recorded for mice in the control group injected with PBS or serum obtained from a rabbit immunized with meningococcal outer membrane preparation. This result clearly indicates that the immunization of another animal species with killed GBS cells can induce the production of antibodies that can passively protect mice. This reagent will also be used to characterize clones.

Table 2 Passive protection of CD-1 mice conferred by rabbit serum obtained after immunization with formaldehyde killed group B whole streptococci (strain C388/90 (Ia/c)) antigenic preparation

groups	number of living mice 14 days after the bacterial challenge with GBS strain C388/90 (Ia/c) ²	% survival
rabbit hyperimmune serum ² - 25 μ l	4/5	80
rabbit hyperimmune serum ¹ - 15 μ l	4/5	80
control rabbit serum - 25 μ l	1/5	20
control PBS	1/10	10

¹ Freund's complete adjuvant was used for first immunization, and Freund's incomplete adjuvant for the following two injections;

² intraperitoneal challenge with 1 ml Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2×10^4 cfu.

EXAMPLE 4 Recombinant production of His.Tag-GBS fusion protein

The coding region of a GBS gene was amplified by PCR (DNA
 5 Thermal Cycler GeneAmp PCR system 2400 Perkin Elmer, San
 Jose, CA) from the genomic DNA of GBS strain C388/90 (Ia/c)
 using the oligos that contained base extensions for the
 addition of the restriction sites BglII (AGATCT) and HindIII
 (AAGCTT), respectively. The PCR product was purified from
 10 agarose gel using a Qiaex II gel extraction kit from Qiagen
 (Chatsworth, CA), digested with the restriction enzymes
 BglII and HindIII (Pharmacia Canada Inc Baie d'Urfe,
 Canada), and extracted with phenol:chloroform before ethanol
 precipitation. The pET-32b(+) vector (Novagen, Madison, WI)
 15 containing the thioredoxin-His.Tag sequence was digested
 with the restriction enzymes BglII and HindIII, extracted
 with phenol:chloroform, and then ethanol precipitated. The
 BglII-HindIII genomic DNA fragment was ligated to the BglII-
 HindIII pET-32b(+) vector to create the coding sequence for
 20 thioredoxin-His.Tag-GBS fusion protein whose gene was under
 control of the T7 promoter. The ligated products were
 transformed into *E. coli* strain XLI Blue MRF' ($\Delta(mcrA)183\Delta$
 (*mcrCB-hsdSMR-mrr*)173 *endA1 supE44 thi-1 recA1 gyrA96 relA1*
lac [F'*proAB lacI*^qZAM15Tn10 (Tet^r)]^c) (Stratagene, La Jolla,
 25 CA) according to the method of Simanis (Hanahan, D. DNA
 Cloning, 1985, D.M. Glover (ed.), pp. 109-135). The
 recombinant pET plasmid was purified using a Qiagen kit
 (Qiagen, Chatsworth, CA) and the nucleotide sequence of the
 DNA insert was verified by DNA sequencing (Taq Dye Deoxy
 30 Terminator Cycle Sequencing kit, ABI, Foster City, CA). The
 recombinant pET plasmid was transformed by electroporation
 (Gene Pulser II apparatus, BIO-RAD Labs, Mississauga,
 Canada) into *E. coli* strain AD494 (DE3) ($\Delta ara^+ leu7697$
 $\Delta lacX74 \Delta phoA PvuII phoR \Delta malF3 F' [lac^+ (lacI^q) pro]$
 35 *trxB::Kan* (DE3)) (Novagen, Madison, WI). In this strain of

E. coli, the T7 promoter controlling expression of the fusion protein, is specifically recognized by the T7 RNA polymerase (present on the λ DE3 prophage) whose gene is under the control of the lac promoter which is inducible by isopropyl- β -D-thio-galactopyranoside (IPTG).

The transformant AD494(DE3)/rpET was grown at 37°C with agitation at 250 rpm in LB broth (peptone 10g/L, Yeast extract 5g/L, NaCl 10g/L) containing 100 μ g of ampicillin (Sigma-Aldrich Canada Ltd., Oakville, Canada) per mL until the A_{600} reached a value of 0.6. In order to induce the production of the thioredoxin-His.Tag-GBS fusion protein, the cells were incubated for 2 additional hours in the presence of IPTG at a final concentration of 1mM. The bacterial cells were harvested by centrifugation.

The recombinant fusion protein produced by AD494(DE3)/rpET32 upon IPTG induction for 2h was partially obtained as insoluble inclusion bodies which were purified from endogenous *E. coli* proteins by the isolation of insoluble aggregates (Gerlach, G.F. et al 1992, Infect. Immun. 60:892). Induced cells from a 500 mL culture were resuspended in 20 mL of 25% sucrose-50mM Tris-HCl buffer (pH8.0) and frozen at -70°C. Lysis of cells in thawed suspension was achieved by the addition of 5mL of a solution of lysozyme (10mg/mL) in 250mM Tris-HCl buffer (pH8.0) followed by an incubation of 10 to 15 min on ice, and the addition of 150mL of detergent mix (5 parts of 20mM Tris-HCl buffer [pH7.4]-300mM NaCl-2% deoxycholic acid-2% Nonidet P-40 and 4 parts of 100mM Tris-HCl buffer [pH8]-50mM EDTA-2% Triton X-100) followed by 5 min incubation on ice. Upon sonication, protein aggregates were harvested by centrifugation for 30 min at 35,000 X g and a sample of the soluble cellular fraction was kept. The aggregated proteins were solubilized in 6M guanidine hydrochloride. The

presence of the fusion protein in both the soluble and insoluble fractions was shown by Western Blot analysis using the serum of a mouse injected with formaldehyde killed cells of GBS strain C388/90 (Ia/c) that survived a bacterial
5 challenge with the corresponding GBS strain.

The purification of the fusion protein from the soluble fraction of IPTG-induced AD494(DE3)/rpET was done by affinity chromatography based on the properties of the
10 His.Tag sequence (6 consecutive histidine residues) to bind to divalent cations (Ni^{2+}) immobilized on the His.Bind metal chelation resin (Novagen, Madison, WI). The purification method used are those described in the pET system Manual, 6th Edition (Novagen, Madison, WI). Briefly, the pelleted
15 cells obtained from a 100mL culture induced with IPTG was resuspended in 4mL of Binding buffer (5mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9), sonicated, and spun at 39,000 X g for 20 min to remove debris. The supernatant was filtered (0.45 μ m pore size membrane) and deposited on a column of
20 His.Bind resin equilibrated in Binding buffer. The column was then washed with 10 column volumes of Binding buffer followed by 6 column volumes of Wash buffer (20mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The thioredoxin-His.Tag-GBS fusion protein was eluted with Elute buffer (1M
25 imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The removal of the salt and imidazole from the sample was done by dialysis against 3 X 1 liter PBS at 4°C.

The quantities of fusion protein obtained from either the
30 soluble or insoluble cytoplasmic fractions of *E. coli* were estimated by Coomassie staining of a sodium dodecyl sulfate (SDS)-polyacrylamide gel with serial dilutions of these proteins and a bovine serum albumin standard (Pierce Chemical Co. Rockford, Ill.).

35

EXAMPLE 5 Recombinant production of GBS protein under
control of lambda P_L promoter

The DNA coding region of a GBS protein was inserted
5 downstream of the promoter λP_L into the translation vector
pURV22. This plasmid was derived from p629 (George et al,
1987, Bio/Technology 5:600) from which the coding region for
a portion of the herpes simplex virus type I (HSV-I)
glycoprotein (gD-1) was removed and the ampicillin
10 resistance gene replaced by a kanamycin cassette obtained
from the plasmid vector pUC4K (Pharmacia Biotech Canada
Inc., Baie D'Urfe, Canada). The vector contained a cassette
of the bacteriophage λ cI857 temperature sensitive repressor
gene from which the functional P_R promoter had been deleted.
15 The inactivation of the cI857 repressor by temperature
increase from the ranges of 30-37°C to 37-42°C resulted in
the induction of the gene under the control of λP_L . The
translation of the gene was controlled by the ribosome
binding site cro followed downstream by a BglII restriction
20 site (AGATCT) and the ATG: ACTAAGGAGGTTAGATCTATG.

Restriction enzymes and T4 DNA ligase were used according to
suppliers (Pharmacia Biotech Canada Inc., Baie D'Urfe,
Canada; and New England Biolabs Ltd., Mississauga, Canada).
25 Agarose gel electrophoresis of DNA fragments was performed
as described by Sambrook et al. (Molecular cloning : A
laboratory Manual, 1989, Cold Spring Harbor Laboratory
Press, N.Y). Chromosomal DNA of the GBS bacteria was
prepared according to procedures described in Jayarao et al
30 (J. Clin. Microbiol., 1991, 29:2774). DNA amplification
reactions by polymerase chain reaction (PCR) were made using
DNA Thermal Cycler GeneAmp PCR system 2400 (Perkin Elmer,
San Jose, CA). Plasmids used for DNA sequencing were
purified using plasmid kits from Qiagen (Chatsworth, CA).
35 DNA fragments were purified from agarose gels using Qiaex II

gel extraction kits from Qiagen (Chatsworth, CA). Plasmid transformations were carried out by the method described by Hanahan (DNA Cloning, Glover (ed.) pp, 109-135, 1985). The sequencing of genomic DNA inserts in plasmids was done using synthetic oligonucleotides which were synthesized by oligonucleotide synthesizer model 394 (the Perkin-Elmer Corp., Applied Biosystems Div. (ABI), Foster City, CA). The sequencing reactions were carried out by PCR using the Taq Dye Deoxy Terminator Cycle Sequencing kit (ABI, Foster City, CA) and DNA electrophoresis was performed on automated DNA sequencer 373A (ABI, Foster City, CA). The assembly of the DNA sequence was performed using the program Sequencer 3.0 (Gene Codes Corporation, Ann Arbor, MI). Analysis of the DNA sequences and their predicted polypeptides was performed with the program Gene Works version 2.45 (Intelligenetics, Inc., Mountain View CA).

The coding region of the GBS gene was amplified by PCR from GBS strain C388/90 (Ia/c) genomic DNA using oligos that contained base extensions for the addition of restriction sites BglII (AGATCT) and XbaI (TCTAGA), respectively. The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and XbaI, and extracted with phenol:chloroform before ethanol precipitation. The pURV22 vector was digested with the restriction enzymes BglII and XbaI, extracted with phenol:chloroform, and ethanol precipitated. The BglII-XbaI genomic DNA fragment was ligated to the BglII-XbaI pURV22 vector in which the GBS gene was under the control of the λ PL promoter. The ligated products were transformed into *E. coli* strain XLI Blue MRF' (Δ (*mcrA*)183 Δ (*mcrCB-hsdSMR-mrr*)173 *endA1 supE44 thi-1 recA1 gyrA96 relA1 lac*[F' *proAB lacI^qZAM15 Tn10(Tet^r)*]^c) (Stratagene, La Jolla CA) according to the methods described in Hanahan, supra. Transformants harboring plasmids with the

insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook et al, supra). The recombinant pURV22 plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of
5 the DNA insert was verified by DNA sequencing.

The transformant XLI Blue MRF'/rpURV22 was grown at 34°C with agitation at 250 rpm in LB broth containing 50µg of kanamycin per mL until the A₆₀₀ reached a value of 0.6. In
10 order to induce the production of the fusion protein, the cells were incubated for 4 additional hours at 39°C. The bacterial cells were harvested by centrifugation , resuspended in sample buffer, boiled for 10 min and kept at -20°C.

15

EXAMPLE 6 Subcloning GBS protein gene in CMV plasmid pCMV-GH

The DNA coding region of a GBS protein was inserted in phase
20 downstream of the human growth hormone (hGH) gene which was under the transcriptional control of the cytomegalovirus (CMV) promoter in the plasmid vector pCMV-GH (Tang et al, Nature, 1992, 356:152). The CMV promoter is non functional in E. coli cells but active upon administration of the
25 plasmid in eukaryotic cells. The vector also incorporated the ampicillin resistance gene.

The coding region of the gene was amplified by PCR from genomic DNA of GBS strain C388/90 (Ia/c) using the oligos
30 that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT). The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII, and
35 extracted with phenol:chloroform before ethanol precipitation. The pCMV-GH vector (Laboratory of Dr. Stephen

A. Johnston, Department of Biochemistry, The University of Texas, Dallas, Texas) containing the human growth hormone to create fusion proteins was digested with the restriction enzymes BamHI and HindIII, extracted with phenol:chloroform, and ethanol precipitated. The 1.3-kb BglII-HindIII genomic DNA fragment was ligated to the BamHI -HindIII pCMV-GH vector to create the hGH-GBS fusion protein under the control of the CMV promoter. The ligated products were transformed into *E. coli* strain DH5 α [ϕ 80 *lacZ* Δ M15 *endA1* *recA1* *hsdR17* (⁺K⁻K⁺) *supE44* *thi-1* λ *gyrA96* *relA1* Δ (*lacZYA-argF*)U169] (Gibco BRL, Gaithersburg, MD) according to the methods described by Hanahan, *supra*. Transformants harboring plasmids with the insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook, J. et al, *supra*). The recombinant pCMV plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

20

EXAMPLE 7 Immunological activity of GBS protein to GBS challenge

Four groups of 12 female CD-1 mice (Charles River, St-Constant, Quebec, Canada) of 6 to 8 weeks were injected subcutaneously three times at three week intervals with 0.1mL of the following antigenic preparations: formaldehyde killed cells of GBS strain C388/90 ($\sim 6 \times 10^7$ cfu), 20 μ g of thioredoxin-His.Tag-GBS fusion protein obtained from the insoluble (inclusion bodies) or 20 μ g of the fusion protein, affinity purified (nickel column), from the soluble cytoplasmic fraction in *E. coli*, or 20 μ g of affinity purified (nickel column) thioredoxin-His.Tag control polypeptide. 20 μ g of QuilATM (Cedarlane Laboratories Ltd, Hornby, Canada)

was added to each antigenic preparation as the adjuvant. Serum samples were obtained from each mouse before immunization (PB) and on days 20 (TB1), 41 (TB2) and 54 (TB3) during the immunization protocols. Sera were frozen at -20°C.

An increase of the ELISA titers was recorded after each injection of the fusion protein indicating a good primary response and a boost of the specific humoral immune response after each of the second and third administration. At the end of the immunization period, the means of reciprocal ELISA titers was 456,145 for the group immunized with 20µg of fusion protein obtained from inclusion bodies compared to 290,133 for the group of mice immunized with the protein from soluble fraction in *E.coli*. The latter result suggests that the protein obtained from inclusion bodies could be more immunogenic than the soluble protein. Analysis of mice sera in ELISA using the affinity purified thioredoxin-His.Tag to coat plates showed that negligible antibody titers are made against the thioredoxin-His.Tag portion of the fusion protein. The reactivity of the sera from mice injected with the recombinant fusion protein was also tested by ELISA against formaldehyde killed whole cells of GBS strain C388/90. The antibodies induced by immunization with recombinant fusion protein also recognized their specific epitopes on GBS cells indicating that their conformation is close enough to the native streptococcal protein to induce cross-reactive antibodies.

To verify whether the immune response induced by immunization could protect against GBS infection, mice were challenged with 3.5×10^5 cfu of GBS strains C338/90(Ia/c) and 1.2×10^5 cfu of strain NCS246(II/R) the results of which are illustrated in tables 3 and 4 respectively. Mice immunized with control thioredoxin-His.Tag peptide were not protected against challenge with either GBS strain while those

immunized with formaldehyde killed C388/90 whole cells only provided protection against homologous challenge. The thioredoxin-His.Tag-GBS fusion protein of the invention protected mice from challenge with both GBS strains. Blood
5 and spleen culture of these mice did not reveal the presence of any GBS.

Table 3 Survival from GBS strain C388/90 (Ia/c) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag ²	1 / 6	17
formaldehyde killed C388/90 cells ³	6 / 6	100
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ⁴	6 / 6	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ⁴	6 / 6	100

- 5 ¹ intraperitoneal administration with 1 ml Todd-Hewitt culture medium adjusted to 3.5×10^5 cfu;
- ² 20 μ g administered; posterior legs paralyzed in surviving mouse; GBS detected in blood and spleen;
- ³ 6×10^7 cfu administered;
- ⁴ 20 μ g administered.

Table 4 Survival from GBS strain NCS246 (II/R) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag ²	0 / 6	0
formaldehyde killed C388/90 cells ³	2 / 6	34
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ²	5 / 5 ⁴	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ²	6 / 6	100

5 ¹ intraperitoneal administration with 1 ml Todd-Hewitt
culture medium containing GBS NCS246 (II/R) suspension
adjusted to 1.2×10^5 cfu.

² 20 µg administered;

³ 6×10^7 cfu administered;

10 ⁴ one mouse died during immunization.

EXAMPLE 8 Immunization with recombinant GBS protein confers protection against experimental GBS infection

15

This example illustrates the protection of mice against
fatal GBS infection by immunization with the recombinant
protein corresponding to the SEQ ID NO:39.

20 Groups of 10 female CD-1 mice (Charles River) were immunized
subcutaneously three times at three-week intervals with 20
µg of recombinant protein purified from E. coli strain BLR
(Novagen) harboring the recombinant pURV22 plasmid vector
containing the GBS gene corresponding to SEQ ID NO:42 in
25 presence of 20 µg of QuilATM adjuvant (Cedarlane
Laboratories Ltd, Hornby, Canada) or, as control, with

QuilA™ adjuvant alone in PBS. Blood samples were collected from the orbital sinus on day 1, 22 and 43 prior to each immunization and fourteen days (day 57) following the third injection. One week later the mice were challenged with approximately 10^4 to 10^6 CFU of various virulent GBS strains. Samples of the GBS challenge inoculum were plated on TSA/5% sheep blood agar plates to determine the CFU and to verify the challenge dose. Deaths were recorded for a period of 14 days and on day 14 post-challenge, the surviving mice were sacrificed and blood and spleen were tested for the presence of GBS organisms. The survival data are shown in table 5.

Prechallenge sera were analyzed for the presence of antibodies reactive with GBS by standard immunoassays. Elisa and immunoblot analyses indicated that immunization with recombinant GBS protein produced in *E. coli* elicited antibodies reactive with both, recombinant and native GBS protein. Antibody responses to GBS are described in Example 9.

20

Table 5. Ability of recombinant GBS protein corresponding to SEQ ID NO: 39 to elicit protection against 8 diverse GBS challenge strains

5

Immunogen	Challenge strain		No. alive: No. dead ¹	
	Designation	Type		
rGBS protein	C388/90	Ia/c	8 : 2	(P<0.0001)
none			0 : 10	
rGBS protein	NCS 246	II/R	10 : 0	(P=0.0012)
none			3 : 7	
rGBS protein	ATCC12401	Ib	10 : 0	(P=0.001)
none			3 : 7	
rGBS protein	NCS 535	V	10 : 0	(P=0.01)
none			5 : 5	
rGBS protein	NCS 9842	VI	10 : 0	(P<0.0001)
none			0 : 10	
rGBS protein	NCS 915	III	7 : 3	(P=0.0007) ²
NCS 915-F ³			1 : 9	
none			4 : 6	
rGBS protein	NCS 954	III/R	7 : 3	(P=0.002)
NCS 954-F			4 : 6	
none			1 : 9	
rGBS protein	COH1	III	4 : 6	(P=0.0004)
COH1-F			3 : 7	
none			0 : 10	

¹ Groups of 10 mice per group were used, the number of mice surviving to infection and the number of dead mice are indicated. The survival curves corresponding to recombinant GBS protein-immunized animals were compared to the survival curves corresponding to mock-immunized animals using the log-rank test for nonparametric analysis.

² Comparison analysis to NCS915-F-immunized animals.

³ Animals were immunized with formaldehyde-killed GBS in presence of QuilATM adjuvant.

All hemocultures from surviving mice were negative at day 14 post-challenge. Spleen cultures from surviving mice were negative except for few mice from experiment MB-11.

EXAMPLE 9 Vaccination with the recombinant GBS protein
elicits an immune response to GBS

Groups of 10 female CD-1 mice were immunized subcutaneously
5 with recombinant GBS protein corresponding to SEQ ID NO:39
as described in Example 8. In order to assess the antibody
response to native GBS protein, sera from blood samples
collected prior each immunization and fourteen days after
the third immunization were tested for antibody reactive
10 with GBS cells by ELISA using plates coated with
formaldehyde-killed GBS cells from type III strain NCS 954,
type Ib strain ATCC12401, type V strain NCS 535 or type VI
strain NCS 9842. The specificity of the raised antibodies
for GBS protein was confirmed by Western blot analyses to
15 GBS cell extracts and purified recombinant antigens. The
results shown in Figure 10 clearly demonstrate that animals
respond strongly to recombinant GBS protein used as
immunogens with median reciprocal antibody titers varying
between 12000 and 128000, for sera collected after the third
20 immunization, depending of the coating antigen. All
preimmune sera were negative when tested at a dilution of
1 :100. GBS-reactive antibodies were detectable in the sera
of each animal after a single injection of recombinant GBS
protein.

25

Example 10 Antigenic conservation of the GBS protein of the present invention

Monoclonal antibodies (MAbs) specific to the GBS protein of the present invention were used to demonstrate that this surface antigen is produced by all GBS and that it is also antigenically highly conserved.

A collection of 68 GBS isolates was used to evaluate the reactivity of the GBS-specific MAbs. These strains were obtained from the National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Canada; Centre Hospitalier Universitaire de Quebec, Pavillon CHUL, Quebec, Canada; American Type Culture Collection, USA; Laboratoire de Sante Publique du Quebec, Canada; and Dept. of Infectious Disease, Children's Hospital and Medical Center, Seattle, USA. All eight MAbs were tested against the following panel of strains: 6 isolates of serotype Ia or Ia/c, 3 isolates of serotype Ib, 4 isolates of serotype II, 14 isolates of serotype III, 2 isolates of serotype IV, 2 isolates of serotype V, 2 isolates of serotype VI, 2 isolates of serotype VII, 1 isolate of serotype VIII, 10 isolates that were not serotyped and 3 bovine *S. agalactiae* strains. MAb 3A2 was also reacted with additional GBS: 9 isolates of serotype Ia/c and 10 isolates of serotype V. The strains were grown overnight on blood agar plates at 37°C in an atmosphere of 5% CO₂. Cultures were stored at -70°C in heart infusion broth with 20% (v/v) glycerol.

To obtain the GBS protein-specific MAbs, mice were immunized three times at three-week intervals with 20 µg of purified recombinant GBS protein (SEQ ID NO :44) in the presence of 20% QuilA™ adjuvant. Hybridoma cell lines were generated by fusion of spleen cells recovered from immunized mice with the nonsecreting SP2/O myeloma cell line as described

previously (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Hybrid clone supernatants were tested for specific antibody production by ELISA using formaldehyde inactivated GBS and purified recombinant GBS protein (SEQ ID NO :39 or 44) as coating antigen, as previously described (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Specific hybrid were cloned by limiting dilutions, expanded, and frozen in liquid nitrogen. Production of recombinant GBS protein was presented in Examples 4 & 5. Purified recombinant GBS protein or formaldehyde inactivated GBS were resolved by electrophoresis by using the discontinuous buffer system of Laemmli as recommended by the manufacturer and then transfer onto nitrocellulose membrane for Western immunoblotting as described previously (Martin et al. 1992. Infect. Immun. 60:2718-2725).

Western immunoblotting experiments clearly indicated that all eight MAbs recognized a protein band that corresponded to the purified recombinant GBS protein (SEQ ID NO :39). These MAbs also reacted with a protein band present in every GBS isolates tested so far. The reactivity of these GBS-specific MAbs are presented in Table 6. Each MAb reacted well with all 46 GBS. In addition, these MAbs also recognized the 3 *S. agalactiae* strains of bovine origin that were tested. MAb 3A2 also recognized nineteen GBS; 9 isolates of serotype Ia/c and 10 of serotype V. The other MAbs were not tested against these additional strains.

These results demonstrated that the GBS protein (SEQ ID NO :39) was produced by all the 65 GBS and the three 3 *S. agalactiae* strains of bovine origin that were tested so far.

More importantly, these results clearly demonstrated that the epitopes recognized by these eight GBS-specific MAbs were widely distributed and conserved among GBS. These results also indicated that these epitopes were not

restricted to serologically related isolates since representatives of all known GBS serotypes including the major disease causing groups were tested.

- 5 In conclusion, the data presented in this example clearly demonstrated that the GBS protein of the present invention is produced by all GBS and that it is antigenically highly conserved.

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Table 6. Reactivity of eight GBS protein-specific MABs with different *S. agalactiae* strains as evaluated by Western immunoblots.

Mabs	Number of each serotype of <i>s. agalactiae</i> strains recognized by the MABs.											
	Ia or Ia/c (6)	Ib (3)	II (4)	III (4)	IV (2)	V (2)	VI (2)	VII (2)	VIII (1)	NT(10) 2	TOTAL (26)	Bovine (3)
3A21	6	3	4	4	2	2	2	2	1	10	46	3
5A12	6	3	4	4	2	2	2	2	1	10	46	3
6G11	6	3	4	4	2	2	2	2	1	10	46	2
8B9	6	3	4	4	2	2	2	2	1	10	46	3
8E11	6	3	4	4	2	2	2	2	1	10	46	3
12B12	6	3	4	4	2	2	2	2	1	10	46	3
18F11	6	3	4	4	2	2	2	2	1	10	46	3
20G2	6	3	4	4	2	2	2	2	1	10	46	3

1 Nine additional strains of serotype Ia/c and 10 strains of serotype V were recognized by MAB 3A2.

10 2 These strains were not serotyped

WE CLAIM:

1. An isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or
fragments, analogs or derivatives thereof.
2. A polynucleotide according to claim 1, wherein said polynucleotide encodes a polypeptide having at least 95% identity to the second polypeptide.
3. An isolated polynucleotide encoding a polypeptide capable of generating antibodies having binding specificity for a polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or
fragments, analogs or derivatives thereof.

4. An isolated polynucleotide that is complementary to the polynucleotide of claim 1.
5. An isolated polynucleotide that is complementary to the polynucleotide of claim 3.
6. The polynucleotide of claim 1, wherein said polynucleotide is DNA.
7. The polynucleotide of claim 3, wherein said polynucleotide is DNA.
8. The polynucleotide of claim 1, wherein said polynucleotide is RNA.
9. The polynucleotide of claim 3, wherein said polynucleotide is RNA.
10. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of :
SEQ ID NO : 1, SEQ ID NO : 7, SEQ ID NO : 13, SEQ ID NO : 22, SEQ ID NO : 27, SEQ ID NO : 32, SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43 or fragments, analogues or derivatives thereof.
11. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of :
SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43.
12. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 37.

13. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 42.
14. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 43.
15. A polynucleotide according to claim 10 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
16. A polynucleotide according to claim 11 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
17. A vector comprising the polynucleotide of claim 1, wherein said polynucleotide is operably linked to an expression control region.
18. A vector comprising the polynucleotide of claim 3, wherein said polynucleotide is operably linked to an expression control region.
19. A host cell transfected with the vector of claim 17.
20. A host cell transfected with the vector of claim 18.
21. A process for producing a polypeptide comprising culturing a host cell according to claim 19 under conditions suitable for expression of said polypeptide.
22. A process for producing a polypeptide comprising culturing a host cell according to claim 20 under condition suitable for expression of said polypeptide.

23. An isolated polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
24. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 39.
25. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 44.
26. An isolated polypeptide capable of generating antibodies having binding specificity for a second polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

27. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO : 39.
28. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO : 44.
29. An isolated polypeptide having an amino acid sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
SEQ ID NO:40 and SEQ ID NO:41 or fragments, analogs or derivatives thereof.
30. The isolated polypeptide of claim 29 having an amino acid sequence according to SEQ ID NO : 39.
31. An isolated polypeptide having an amino acid sequence according to SEQ ID NO : 44.
32. An isolated polypeptide according to any one of claims 29 to 31, wherein the N-terminal Met residue is deleted.
33. An isolated polypeptide according to any one of claims 29 to 30, wherein the secretory amino acid sequence is deleted.
34. A vaccine composition comprising a polypeptide according to any one of claims 23 to 31 and a pharmaceutically acceptable carrier, diluent or adjuvant.

35. A vaccine composition comprising a polypeptide according to claim 32 and a pharmaceutically acceptable carrier, diluent or adjuvant.
36. A vaccine composition comprising a polypeptide according to claim 33 and a pharmaceutically acceptable carrier, diluent or adjuvant.
37. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 34.
38. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 35.
39. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 36.
40. A method according to any one of claims 37 to 39, wherein said animal is a bovine.
41. A method according to any one of claims 37 to 39, wherein said animal is a human.

42. A method according to any one of claims 37 to 39, wherein said bacterial infection is selected from the group consisting of group A streptococcus and group B streptococcus.
43. A method according to claim 42, wherein said bacterial infection is group B streptococcus.
44. Use of a vaccine composition according to claim 34 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
45. Use of a vaccine composition according to any one of claims 35 to 36 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
46. Use of a vaccine composition according to any one claims 23 to 31 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
47. Use of a vaccine composition according to claim 32 for the manufacture of a vaccine for the therapeutic or

prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

48. Use of a vaccine composition according to claim 33 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

TATCTGGCAA AGAGCCAGCT AATCGTTTGA GTTGGGCTAA AAATAAATTA TTAATCAATG 60
 S G K E P A N R F S W A K N K L L I N G
 ---->
 GATTCAATGC AACTCTAGCA GCAACTATCT TATTTTTTGC AGTTCAATTC ATAGGTCTTA 120
 F I A T L A A T I L F F A V Q F I G L K
 AACCAGATTA CCCTGGAAAA ACCTACTTTA TTATCTTATT GACAGCATGG ACTTTGATGG 180
 P D Y P G K T Y F I I L L T A W T L M A
 CATTAGTAAC TGCTTTAGTG GGATGGGATA ATAGGTATGG TTCCTTCTTG TCGTTATTAA 240
 L V T A L V G W D N R Y G S F L S L L I
 TATTATTATT CCAGCTTGGT TCAAGCGCAG GAACTTACCC AATAGAATTG AGTCCTAAGT 300
 L L F Q L G S S A G T Y P I E L S P K F
 TCTTCAAC AATTCAACCA TTTTACC GA TACTACTC TGTTTCAGGA TTAAGAGAGA 360
 F Q T I Q P F L P M T Y S V S G L R E T
 CCATCTCGTT GACGGGAGAC GTTAACCATC AATGGAGAAT GCTAGTAATC TTTTATGAT 420
 I S L T G D V N H Q W R M L V I F L V S
 CATCGATGAT ACTTGCTCTT CTTATTTATC GTAAACAAGA AGATTAAATAG AAAGTATCTA 480
 S M I L A L L I Y R K Q E D
 GTGATAGACT AACAGTATGA TATGGTATGT CAAAGTATTT AGGAGGAGAA GATATGTCTA 540
 M S T
 |---->
 CTTTAACAAT AATTATTGCA ACATTAACCTG CTTTGAACA TTTTATATT ATGTATTGG 600
 L T I I I A T L T A L E H F Y I M Y L E
 AGACGTTAGC CACCCAGTCA AATATGACTG GGAAGATTTT TAGTATGTCT AAAGAAGAGT 660
 T L A T Q S N M T G K I F S M S K E E L
 TGTCAATATTT ACCCGTTATT AAACCTTTTA AGAATCAAGG TGTATACAAC GGCTTGATTG 720
 S Y L P V I K L F K N Q G V Y N G L I G
 GCCTATTCCT CCTTATGGG TTATATATTT CACAGAATCA AGAAATTGTA GCTGTTTTTT 780
 L F L L Y G L Y I S Q N Q E I V A V F L
 TAATCAATGT ATGCTAGTT GCTATTTATG GTGCTTTGAC AGTTGATAAA AAAATCTTAT 840
 I N V L L V A I Y G A L T V D K K I L L
 TAAAACAGGG TGGTTTACCT ATATTAGCTC TTTTAAACATT CTTATTTTAA TACTACTTAG 900
 K Q G G L P I L A L L T F L F
 CCGTTCGATT TAGTTGAACG GCTTTTAGTA ATCATTTTTT TCTCATAATA CAGGTAGTTT 960
 AAGTAATTTG TCTTTAAAA TAGTATAATA TAACTACGAA TTCAAAGAGA GGTGACTTTG 1020
 ATTATGACTG AGAACTGGTT ACATACTAAA GATGGTTCAG ATATTTATTA TCGTGTCTGT 1080
 M T E N W L H T K D G S D I Y Y R V V
 |---->
 GGTCAAGGTC AACCGATTGT TTTTATCAT GGCAATAGCT TAAGTAGTCG CTATTTTGAT 1140
 G Q G Q P I V F L H G N S L S S R Y F D
 AAGCAAATAG CATATTTTTC TAAGTATTAC CAAGTTATTG TTATGGATAG TAGAGGGCAT 1200
 K Q I A Y F S K Y Y Q V I V M D S R G H
 GGCAAAAGTC ATGCAAAGCT AAATACCATT AGTTTCAGGC AAATAGCAGT TGAATTAAG 1260
 G K S H A K L N T I S F R Q I A V D L K

GATATCTTAG TTCATTTAGA GATTGATAAA GTTATATTGG TAGGCCATAG CGATGGTGCC 1320
 D I L V H L E I D K V I L V G H S D G A
 AATTTAGCTT TAGTTTTTCA AACGATGTTT CCAGGTATGG TTAGAGGGCT TTTGCTTAAT 1380
 N L A L V F Q T M F P G M V R G L L L N
 TCAGGGAACC TGA CTATTCA TGGTCAGCGA TGGTGGGATA TTCTTTTAGT AAGGATTGCC 1440
 S G N L T I H G Q R W W D I L L V R I A
 TATAAATTC TTTACTATTT AGGGAACTC TTTCCGTATA TGAGGCAAAA AGCTCAAGTT 1500
 Y K F L H Y L G K L F P Y M R Q K A Q V
 ATTTGCTTA TGTGGAGGA TTTGAAGATT AGTCCAGCTG ATTTACAGCA TGTGTCAACT 1560
 I S L M L E D L K I S P A D L Q H V S T
 CCTGTAATGG TTTTGGTTGG AAATAAGGAC ATAATTAAGT TAAATCATTC TAAGAACTT 1620
 P V M V L V G N K D I I K L N H S K K L
 GCTTCTTATT TTCCAAGGGG GGAGTTTAT TCTTTAGTTG GCTTTGGGCA TCACATTATT 1680
 A S Y F P R G E F Y S L V G F G H H I I
 AAGCAAGATT CCCATGTTTT TAATATTATT GCAAAAAAGT TTATCAACGA TACGTTGAAA 1740
 K Q D S H V F N I I A K K F I N D T L K
 GGAGAAATTG TTGAAAAGC TAATTGAAAA AGTCAAATCA CTGACTTCTG TGATTAAAT 1800
 G E I V E K A N
 TGTATTTTTT ATATCTGTTT TAGTGCTTAT TATTGTTGAA ATGATTCATT TGAAACGAAC 1860
 M I H L K R T
 |---->
 TATTTCTGTT GAGCACTAA AGAGTGT TGGGCAATTA TCTCCAATGA ATCTTTTCTT 1920
 I S V E Q L K S V F G Q L S P M N L F L
 AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGATATGACT TTGTACTGAA 1980
 I I L V G V I A V L P T T G Y D F V L N
 TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTTA CAGACTAGTT GGTGTATCAA 2040
 G L L R T D K S K R Y I L Q T S W C I N
 CACTTTTAAT AACTTGTCAG GATTCGGTGG CTTAATCGAT ATTGGGTTGC GCATGGCTTT 2100
 T F N N L S G F G G L I D I G L R M A F
 TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTGACTCGTT TTTTACCCTA 2160
 Y G K K G Q E K S D L R E V T R F L P Y
 TCTTATTTCT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATCATGAGCC ATATTTTCA 2220
 L I S G L S F I S V I A L I M S H I F H
 TGCCAAAGCT AGTGTGATT ACTATTATTT GGTATTAATT GGTGCTAGTA TGTATTTTCC 2280
 A K A S V D Y Y Y L V L I G A S M Y F P
 TGTATTTAT TGGATTTCTG GTCATAAAGG AAGCCATTAT TTCGGAGATA TGCCATCTAG 2340
 V I Y W I S G H K G S H Y F G D M P S S
 TACTCGTATA AAATTAGGTG TTGTTTCTTT TTTTGAATGG GGATGTGCGG CCGCAGCATT 2400
 T R I K L G V V S F F E W G C A A A A F
 TATAATTATC GGTTATTTAA TGGGCATTCA TCTACCAGTT TATAAAATTT TACCACTATT 2460
 I I I G Y L M G I H L P V Y K I L P L F

TTGTATTGGT TGTGCCGTCG GGATTGTATC CCTTATTCCC GGTGGATTAG GAAGTTTGA 2520
 C I G C A V G I V S L I P G G L G S F E
 ATTAGTTCTA TTTACAGGGT TTGCTGCCGA GGGACTACCT AAAGAACTG TGGTTGCATG 2580
 L V L F T G F A A E G L P K E T V V A W
 GTTATTACTT TATCGTTTAG CCTACTATAT TATTCCATTC TTTGCAGGTA TCTATTTCTT 2640
 L L L Y R L A Y Y I I P F F A G I Y F F
 TATCCATTAT TTAGGTAGTC AAATAAATCA ACGTTATGAA AATGTCCCGA AAGAGTTAGT 2700
 I H Y L G S Q I N Q R Y E N V P K E L V
 ATCAACTGTT CTACAAACCA TGGTGAGCCA TTTGATGCGT ATTTTAGGTG CATTCTTAAT 2760
 S T V L Q T M V S H L M R I L G A F L I
 |---->
 ATTTTCAACA GCATTTTTTG AAAATATTAC TTATATTATG TGGTTGCAGA AGCTAGGCTT 2820
 F S T A F F E N I T Y I M W L Q K L G L
 GGACCCATTA CAAGAACAAA TGTTATGGCA GTTTCAGGT TTATTGCTGG GGGTTTGT 2880
 D P L Q E Q M L W Q F P G L L L G V C F
 TATTCTCTTA GCTAGAACTA TTGATCAAAA AGTGAAAAAT GCTTTTCCAA TTGCTATTAT 2940
 I L L A R T I D Q K V K N A F P I A I I
 CTGGATTACT TTGACATTGT TTTATCTTAA TTTAGGTCAT ATTAGTTGGC GACTATCTTT 3000
 W I T L T L F Y L N L G H I S W R L S F
 CTGGTTTATT TTAATTATTGT TAGGCTTATT AGTCATTAAG CCAACTCTCT ATAAAAACA 3060
 W F I L L L L G L L V I K P T L Y K K Q
 ATTTATTAT AGCTGGGAAG AGCGTATTAA GGATGGAATC ATTATCGTTA GTTTAATGGG 3120
 F I Y S W E E R I K D G I I I V S L M G
 AGTTCTATTT TATATTGCAG GACTACTATT CCCTATCAGG GCTCATATTA CAGGTGGTAG 3180
 V L F Y I A G L L F P I R A H I T G G S
 TATTGAACGC CTGCATTATA TCATAGCATG GGAGCCGATA GCATTGGCTA CGTTGATTCT 3240
 I E R L H Y I I A W E P I A L A T L I L
 TACTCTCGTT TATTTATGTT TGGTTAAGAT TTTACAAGGA AAATCTTGTC AGATTGGTGA 3300
 T L V Y L C L V K I L Q G K S C Q I G D
 TGTGTTCAAT GTGGATCGTT ATAAAAAACT ACTTCAAGCT TACGGTGGTT CTTCGGATAG 3360
 V F N V D R Y K K L L Q A Y G G S S D S
 CGGTTTAGCC TTTTAAATG ATAAAAGGCT CTAAGGTAC CAAAAAATG GAGAAGATTG 3420
 G L A F L N D K R L Y W Y Q K N G E D C
 CGTTGCGTTC CAATTGTAA TTGTCAATAA TAAATGTCTT ATTATGGGGG AACCAGCCGG 3480
 V A F Q F V I V N N K C L I M G E P A G
 TGATGACACT TATATTCGTG AAGCTATTGA ATCGTTTATT GATGATGCTG ATAAGCTAGA 3540
 D D T Y I R E A I E S F I D D A D K L D
 CTATGACCTT GTTTTTTACA GTATTGGACA GAAGTTGACA CTACTTTTAC ATGAGTATGG 3600
 Y D L V F Y S I G Q K L T L L L H E Y G
 TTTTGACTTT ATGAAAGTTG GTGAGGATGC TTTAGTTAAT TTAGAAACGT TTACTCTTAA 3660
 F D F M K V G E D A L V N L E T F T L K

AGGGAATAAG TACAAACCTT TCAGAAATGC CCTAAATAGA GTTGAAAAGG ATGGTTTCTA 3720
G N K Y K P F R N A L N R V E K D G F Y

TTTCGAAGTT GTACAATCGC CACATAGTCA AGAGCTACTA AATAGTTTGG AAGAGATTTC 3780
F E V V Q S P H S Q E L L N S L E E I S

TAATACTTGG TTAGAAGGAC GTCCTGAAAA AGGTTTCTCA CTAGGATATT TTAATAAAGA 3840
N T W L E G R P E K G F S L G Y F N K D

TTATTTCCAA CAAGCCCCAA TAGCTTTGGT AAAAAATGCT GAACACGAAG TTGTTGCTTT 3900
Y F Q Q A P I A L V K N A E H E V V A F

TGCTAATATT ATGCCAAACT ATGAAAAGAG TATTATCTCT ATTGATTTAA TGCGTCACGA 3960
A N I M P N Y E K S I I S I D L M R H D

TAAACAGAAA ATTCCGAATG GCGTTATGGA TTTCTCTTTT TTATCATTAT TCTCTTATTA 4020
K Q K I P N G V M D F L F L S L F S Y Y

TCAAGAGAAG GGATACCACT ATTTTGATT GGGGATGGCA CCTTTATCAG GAGTTGGTCG 4080
Q E K G Y H Y F D L G M A P L S G V G R

CGTTGAAACA AGTTTGTCTA AAGAGAGAAT GCGTATCTT GTCTATCATT TCGGTAGTCA 4140
V E T S F A K E R M A Y L V Y H F G S H

TTTCTACTCA TTTAATGGTT TACACAAGTA TAAGAAGAAG TTTACACCAT TGTGGTCGGA 4200
F Y S F N G L H K Y K K K F T P L W S E

ACGTTATATT TCTTGTCTC GTTCGTCCTG GTTAATTGT GCTATTGTG CCCTATTAAT 4260
R Y I S C S R S S W L I C A I C A L L M

GGAAGATAGT AAAATTAAGA TTGTTAAATA AGCTTTATTT GGCAATTAAA AAGAGCATGT 4320
E D S K I K I V K

CATGCGACAT GCTCTTTTAA AATCATTTAA TACCATTGAT TGCTTGAATC TACTTTATAA 4380

TATGATGTGC TTTTAAATAT TGTTTAGCTA CTGTAGCTGC TGATTATATG TTTACAGCTA 4440

CTTGGTAGTT CATTTCTTGC ATTTCTTTTT CAGTGATATG ACCAGCAAGT TTATTGAGAG 4500

CTTTTTTTTAC TTGA (SEQ ID NO:1) 4514

FIG. 1a
[clone1-dna/aa]

SGKEPANRFS WAKNKLLING FIATLAATIL FFAVQFIGLK PDYPGKTYFI 50
ILLTAWTLMA LVTALVGWDN RYGSFSLSLI LLFQLGSSAG TYPIELSPKF 100
FQTIQPF LPM TYSVSG LRET ISLTGDVNHQ WRMLVIFLVS SMILALLIYR 150
KQED (SEQ ID NO:2) 154

FIG. 1b

MSTLTIIIIAT LTALEHFIYIM YLETLATQSN MTGKIFSMSK EELSYLPVIK 50
LFTNQGVYNG LIGLFLLYGL YISQSQEIVA VFLINVLLVA IYGALTVDKK 100
ILLKQGG LPI LALLTFLF (SEQ ID NO:3) 118

FIG. 1c

MTENWLHTKD GSDIYYRVVG QGQPIVFLHG NSLSSRYFDK QIAYFSKYYQ 50
VIVMDSRGHG KSHAKLNTIS FRQIAVDLKD ILVHLEIDKV ILVGHS DGAN 100
LALVFQTMFP GMVRGLLLNS GNLTIHGQRW WDILLVRIAY KFLHYLGKLF 150
PYMRQKAQVI SLMLDLKIS PADLQHVSTP VMVLVG NKDI IKLNHSSKLA 200
SYFPRGEFYS LVGFGHHIHK QDSHVFNIIA KKFINDTLKG EIVEKAN 247
(SEQ ID NO:4)

FIG. 1d

MIHLKRTISV	EQLKSVFGQL	SPMNLFLIIL	VGVI AVLPTT	GYDFVLNGLL	50
RTDKSKRYIL	QTSWCINTFN	NLSGFGGLID	IGLRMAFYGK	KGQEKSDLRE	100
VTRFLPYLIS	GLSFISVIAL	IMSHIFHAKA	SVDYYYLVLI	GASMYFPVIY	150
WISGHKGSHY	FGDMPSSTRI	KLGVSFFFEW	GCAAAAFIII	GYLMGIHLPV	200
YKILPLFCIG	CAVGIVSLIP	GGLGSFELVL	FTGF AAEGLP	KETVVAWLLL	250
YRLAYYIIPF	FAGIYFFIHY	LGSQINQRYE	NVPKELVSTV	LQTMVSHLMR	300
ILGAFLIFST	AFFENITYIM	WLQKLGLDPL	QEQLMWQFPG	LLLGVCFILL	350
ARTIDQKVKN	AFPIAIIWIT	LTLFYLN LGH	ISWRLSFWFI	LLLLGLLVIK	400
PTLYKKQFIY	SWEERIKDGI	IIVSLMGVLF	YIAGLLFPPIR	AHITGGSIER	450
LHYIIAWEPI	ALATLILTLV	YLCLVKILQG	KSCQIGDVFN	VDRYKKLLQA	500
YGGSSSDSLA	FLNDKRLYWY	QKNGEDCVAF	QFVIVNNKCL	IMGEPAGDDT	550
YIREAIESFI	DDADKLDYDL	VFYSIGQKLT	LLLHEYGFDF	MKVGEDALVN	600
LETFTLKG NK	YKPF RNALNR	VEKDGFYFEV	VQSPHSQELL	NSLEEISNTW	650
LEGRPEKGFS	LGYFNKDYFQ	QAPIALVKNA	EHEVVAFANI	MPNYEKSIIS	700
IDLMRHDQKQ	IPNGVMDFLF	LSLFSYYQEK	GYHYFDLGMA	PLSGVGRVET	750
SFAKERMAYL	VYHFGSHFYS	FNLHKKYKKK	FTPLWSERYI	SCSRSSWLIC	800
AICALLMEDS	KIKIVK	(SEQ ID NO:5)			816

FIG. 1e

MRILGAFLIF	STAFFENITY	IMWLQKLGLD	PLQEQLMWQF	PGLLLGVCFI	50
LLARTIDQKV	KNAFPIAIIW	ITLTLFYLN L	GHISWRLSFW	FILLLLGLLV	100
IKPTLYKKQF	IYSWEERIKD	GIIIVSLMGV	LFYIAGLLFP	IRAHITGGS I	150
ERLHYIIAWE	PIALATLILT	LVYLCLVKIL	QGKSCQIGDV	FNVDRYKKLL	200
QAYGGSSDSG	LAFLNDKRLY	WYQKNGEDCV	AFQFVIVNNK	CLIMGEPAGD	250
DTYIREAIES	FIDDADKLDY	DLVFYSIGQK	LTLLHEYGF	DFMKVGEDAL	300
VNLETFTLKG	NKYKPF RNAL	NRVEKDGFYF	EVVQSPHSQE	LLNSLEEISN	350
TWLEGRPEKG	FSLGYFNKDY	FQQAPIALVK	NAEHEVVAF A	NIMPNYEKSI	400
ISIDLMRHDK	QKIPNGVMDF	LFLSLFSYYQ	EKGYHYFDLG	MAPLSGVGRV	450
ETSFAKERMA	YLVYHFGSHF	YSFNLHKKYK	KKFTPLWSER	YISCSRSSWL	500
ICAICALLME	DSKIKIVK	(SEQ ID NO:6)			518

FIG. 1f

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AATTTTGATA TCGAAACAAC AACTTTTGAG GCAATGAAAA AGCACGCGTC ATTATTGGAG    60
N F D I E T T T F E A M K K H A S L L E
----->
AAAATATCTG TTGAGCGTTC TTTTATTGAA TTTGATAAAC TTCTATTAGC ACCTTATTGG    120
K I S V E R S F I E F D K L L L A P Y W

CGTAAAGGAA TGCTGGCACT AATAGATAGT CATGCTTTTA ATTATCTACC ATGCTTAAAA    180
R K G M L A L I D S H A F N Y L P C L K

AATAGGGAAT TACAATTAAG CGCCTTTTGT TCCCAGTTAG ATAAAGATTT TTTATTGAG    240
N R E L Q L S A F L S Q L D K D F L F E

ACATCAGAAC AAGCTTGGGC ATCACTCATC TTGAGTATGG AAGTTGAACA CACAAAGACT    300
T S E Q A W A S L I L S M E V E H T K T

TTTTTAAAAA AATGGAAGAC ATCAACTCAC TTCAAAAAG ATGTTGAGCA TATAGTGGAT    360
F L K K W K T S T H F Q K D V E H I V D

GTTTATCGTA TTCGTGAACA AATGGGATTG GCTAAAGAAC ATCTTTATCG TTATGGAAAA    420
V Y R I R E Q M G L A K E H L Y R Y G K

ACTATAATAA AACAAGCGGA AGGTATTCGC AAAGCAAGAG GCTTGATGGT TGATTTCGAA    480
T I I K Q A E G I R K A R G L M V D F E

AAAATAGAAC AACTAGATAG TGAGTTAGCA ATCCATGATA GGCATGAGAT AGTTGTCAAT    540
K I E Q L D S E L A I H D R H E I V V N

GGTGGCACCT TAATCAAGAA ATTAGGAATA AAACCTGGTC CACAGATGGG AGATATTATC    600
G G T L I K K L G I K P G P Q M G D I I

TCTCAAATTG AATTAGCCAT TGTTTATAGG CAACTGATTA ATGAAGAAGA GGCTATTTTA    660
S Q I E L A I V L G Q L I N E E E A I L

CATTTTGTTA AGCAGTACTT GATGGATTAG AGAGGATTAT ATGAGCGATT TTTAGTAGA    720
H F V K Q Y L M D M S D F L V D
|----->
TGGATTGACT AAGTCGGTTG GTGATAAGAC GGTCTTTAGT AATGTTTCAT TTATCATCCA    780
G L T K S V G D K T V F S N V S F I I H

TAGTTTAGAC CGTATTGGGA TTATTGGTGT CAATGGAAC TGGAAAGACAA CACTATTAGA    840
S L D R I G I I G V N G T G K T T L L D

TGTTATTTTCG GGTGAATTAG GTTTTGATGG TGATCGTTCC CCTTTTTCAT CAGCTAATGA    900
V I S G E L G F D G D R S P F S S A N D

TTATAAGATT GCTTATTTAA AACAAGAACC AGACTTTGAT GATTCTCAGA CAATTTTGGA    960
Y K I A Y L K Q E P D F D D S Q T I L D

CACCGTACTT TCTTCTGACT TAAGAGAGAT GGCTTTAATT AAAGAATATG AATTATTGCT    1020
T V L S S D L R E M A L I K E Y E L L L

TAATCACTAC GAAGAAAGTA AGCAATCACG TCTAGAGAAA GTAATGGCAG AAATGGATTC    1080
N H Y E E S K Q S R L E K V M A E M D S

TTTAGATGCT TGGTCTATTG AGAGCGAAGT CAAAACAGTA TTATCCAAAT TAGGTATTAC    1140
L D A W S I E S E V K T V L S K L G I T

TGATTTGCAG TTGTCGGTTG GTGAATTATC AGGAGGATTA CGAAGACGTG TTCAATTAGC    1200
D L Q L S V G E L S G G L R R R V Q L A

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GCAAGTATTA	TTAAATGATG	CAGATTTATT	GCTCTTAGAC	GAACCTACTA	ACCACTTAGA	1260
Q V L	L N D A	D L L	L L D	E P T N	H L D	
TATTGACACT	ATTGCATGGT	TAACGAATTT	TTTGAAAAAT	AGTAAAAAGA	CAGTGCTTTT	1320
I D T	I A W L	T N F	L K N	S K K T	V L F	
TATAACTCAT	GATCGTTATT	TTCTAGACAA	TGTTGCAACA	CGTATTTTGG	AATTAGATAA	1380
I T H	D R Y F	L D N	V A T	R I F E	L D K	
GGCACAGATT	ACAGAATATC	AAGGCAATTA	TCAGGATTAT	GTCCGACTTC	GTGCAGAACA	1440
A Q I	T E Y Q	G N Y	Q D Y	V R L R	A E Q	
AGACGAGCGT	GATGCTGCTA	GTTTACATAA	AAAGAAACAG	CTTATAAAC	AGGAAC TAGC	1500
D E R	D A A S	L H K	K K Q	L Y K Q	E L A	
TTGGATGCGT	ACTCAGCCAC	AAGCTCGTGC	AACGAAACAA	CAGGCTCGTA	TTAATCGTTT	1560
W M R	T Q P Q	A R A	T K Q	Q A R I	N R F	
TCAAAATCTA	AAAAACGATT	TACACCAAAC	AAGCGATACA	AGCGATTTGG	AAATGACATT	1620
Q N L	K N D L	H Q T	S D T	S D L E	M T F	
TGAAACAAGT	CGAATTGGGA	AAAAGGTTAT	TAATTTTGAA	AATGTCTCTT	TTTCTTACCC	1680
E T S	R I G K	K V I	N F E	N V S F	S Y P	
AGATAAATCT	ATCTTGAAAG	ACTTTAATTT	GTTAATTCAA	AATAAGACC	GTATTGGCAT	1740
D K S	I L K D	F N L	L I Q	N K D R	I G I	
CGTTGGAGAT	AATGGTGTG	GAAAGTCAAC	CTTACTTAAT	TTAATTGTTT	AAGATTTACA	1800
V G D	N G V G	K S T	L L N	L I V Q	D L Q	
GCCGGATTCTG	GGTAATGTCT	CTATTGGTGA	AACGATACGT	GTAGGTTACT	TTTCACAACA	1860
P D S	G N V S	I G E	T I R	V G Y F	S Q Q	
ACTTCATAAT	ATGGATGGCT	CAAAACGTGT	TATTAATTAT	TTGCAAGAGG	TTGCAGATGA	1920
L H N	M D G S	K R V	I N Y	L Q E V	A D E	
GGTTAAAACT	AGTGTGCGTA	CAACAAGTGT	GACAGAACTA	TTGGAACAAT	TTCTCTTTCC	1980
V K T	S V G T	T S V	T E L	L E Q F	L F P	
ACGTTTCGACA	CATGGAACAC	AAATTGCAAA	ATTATCAGGT	GGTGAGAAAA	AAAGACTTTA	2040
R S T	H G T Q	I A K	L S G	G E K K	R L Y	
CCTTTTAAAA	ATCCTGATTG	AAAAGCCTAA	TGTGTTACTA	CTTGATGAGC	CGACAAATGA	2100
L L K	I L I E	K P N	V L L	L D E P	T N D	
CTTAGATATT	GCTACATTAA	CTGTTCTTGA	AAATTTTTTA	CAAGGCTTTG	GTGGTCCTGT	2160
L D I	A T L T	V L E	N F L	Q G F G	G P V	
GATTACAGTT	AGTCACGATC	GTTACTTTTT	AGATAAAGTG	GCTAATAAAA	TTATTGCGTT	2220
I T V	S H D R	Y F L	D K V	A N K I	I A F	
TGAAGATAAC	GATATCCGTG	AATTTTTTGG	TAATTATACT	GATTATTTAG	ATGAAAAAGC	2280
E D N	D I R E	F F G	N Y T	D Y L D	E K A	
ATTTAATGAG	CAAAATAATG	AAGTTATCAG	TAAAAAAGAG	AGTACCAAGA	CAAGTCGTGA	2340
F N E	Q N N E	V I S	K K E	S T K T	S R E	
AAAGCAAAGT	CGTAAAAGAA	TGTCTTACTT	TGAAAAACAA	GAATGGGCGA	CAATTGAAGA	2400
K Q S	R K R M	S Y F	E K Q	E W A T	I E D	
CGATATTATG	ATATTGGAAG	ATACTATCAC	TCGTATAGAA	AATGATATGC	AAACATGTGG	2460

D I M I L E N T I T R I E N D M Q T C G
 TAGTGATTTT ACAAGGTTAT CTGATTTACA AAAGGAATTA GATGCAAAAA ATGAAGCACT 2520
 S D F T R L S D L Q K E L D A K N E A L
 TCTAGAAAAG TATGACCGTT ATGAGTACCT TAGTGAGTTA GACACATGAT TATCCGTCCG 2580
 L E K Y D R Y E Y L S E L D T M I I R P
 |----->
 ATTATTAAAA ATGATGACCA AGCAGTTGCA CAATTAATTC GACAAAGTTT ACGCGCCTAT 2640
 I I K N D D Q A V A Q L I R Q S L R A Y
 GATTTAGATA AACCTGATAC AGCATATTCA GACCCTCACT TAGATCATTT GACCTCATAC 2700
 D L D K P D T A Y S D P H L D H L T S Y
 TACGAAAAAA TAGAGAAGTC AGGATTCTTT GTCATTGAGG AGAGAGATGA GATTATTGGC 2760
 Y E K I E K S G F F V I E E R D E I I G
 TGTGGCGGCT TTGGTCCGCT GAAAAATCTA ATTGCAGAGA TGCAGAAGGT GTACATTGCA 2820
 C G G F G P L K N L I A E M Q K V Y I A
 GAACGTTTCC GTGGTAAGGG GCTTGCTACT GATTTAGTGA AAATGATTGA AGTAGAAGCT 2880
 E R F R G K G L A T D L V K M I E V E A
 CGAAAAATTG GGTATAGACA ACTTTATTTA GAGACAGCCA GTACTTTGAG TAGGGCAACT 2940
 R K I G Y R Q L Y L E T A S T L S R A T
 GCGGTTTATA AGCATATGGG ATATTGTGCC TTATCGCAAC CAATAGCAAA TGATCAAGGT 3000
 A V Y K H M G Y C A L S Q P I A N D Q G
 CATACAGCTA TGGATATTTG GATGATTAAA GATTATAAG TTGAAAGTGG ATTAGTGAAC 3060
 H T A M D I W M I K D L
 ATGGATTAAT TATTTTGAGA TAAGAGGAAA GAAAAGGAGA CATATATGGC ATATATTTGG 3120
 M A Y I W
 |----->
 TCTTATTGA AAAGGTACCC CAATTGGTTA TGGCTTGATT TACTAGGAGC TATGCTTTTT 3180
 S Y L K R Y P N W L W L D L L G A M L F
 GTGACGGTTA TCCTAGGAAT GCCACAGCC TTAGCGGGTA TGATTGATAA TGGCGTTACA 3240
 V T V I L G M P T A L A G M I D N G V T
 AAAGGTGATC GGAAGTGGAGT TTATCTGTGG ACGTTCATCA TGTTTATATT TGTTGTACTA 3300
 K G D R T G V Y L W T F I M F I F V V L
 GGTATTATTG GGCCTATTAC GATGGCTTAC GCATCTAGTC GCTTAACGAC AACAAATGATT 3360
 G I I G R I T M A Y A S S R L T T T M I
 AGAGATATGC GTAATGATAT GTATGCTAAG CTTCAAGAAT ACTCCCATCA TGAATATGAA 3420
 R D M R N D M Y A K L Q E Y S H H E Y E
 CAGATAGGTG TATCTTCACT AGTGACACGT ATGACAAGCG ATACTTTTGT TTTGATGCAA 3480
 Q I G V S S L V T R M T S D T F V L M Q
 TTTGCTGAAA TGTCTTTACG TTTAGGCCTA GTAACCTCTA TGGTAATGAT TTTTAGCGTG 3540
 F A E M S L R L G L V T P M V M I F S V
 GTTATGATAC TAATTACGAG TCCATCTTTG GCTTGGCTTG TAGCGGTTGC GATGCCTCTT 3600
 V M I L I T S P S L A W L V A V A M P L
 TTGGTAGGAG TCGTTTTATA TGTAGCTATA AAAACAAAAC CTTTATCTGA AAGACAACAG 3660
 L V G V V L Y V A I K T K P L S E R Q Q

ACTATGCTTG ATAAAATCAA TCAATATGTT CGTGAAAATT TAACAGGGTT ACGCGTTGTT 3720
T M L D K I N Q Y V R E N L T G L R V V

AGAGCCTTTG CAAGAGAGAA TTTTCAATCA CAAAAATTC AAGTCGCTAA CCAACGTTAC 3780
R A F A R E N F Q S Q K F Q V A N Q R Y

ACAGATACTT CAACTGGTCT TTTTAAATTA ACAGGGCTAA CAGAACCACT TTTCGTTCAA 3840
T D T S T G L F K L T G L T E P L F V Q

ATTATTATTG CAATGATTGT GGCTATCGTT TGGTTTGCTT TGGATCCCTT ACAAAGAGGT 3900
I I I A M I V A I V W F A L D P L Q R G

GCTATTAAAA TAGGGGATTT AGTTGCTTTT ATCGAATATA GCTTCCATGC TCTCTTTTCA 3960
A I K I G D L V A F I E Y S F H A L F S

TTTTTGCTAT TTGCCAATCT TTTTACTATG TATCCTCGTA TGGTGGTATC AAGCCATCGT 4020
F L L F A N L F T M Y P R M V V S S H R

ATTAGAGAGG TGATGGATAT GCCAATCTCT ATCAATCCTA ATGCCGAAGG TGTTACGGAT 4080
I R E V M D M P I S I N P N A E G V T D

ACGAACTTA AAGGGCATT AGAATTTGAT AATGTAACAT TCGCTTATCC AGGAGAAACA 4140
T K L K G H L E F D N V T F A Y P G E T

GAGAGTCCCG TTTTGCATGA TATTTCTTTT AAAGCTAAGC CTGGAGAAAC AATTGCTTTT 4200
E S P V L H D I S F K A K P G E T I A F

ATTGGTTCAA CAGGTTCAAG AAAATCTTCT CTTGTTAATT TGATTCCACG TTTTATGAT 4260
I G S T G S G K S S L V N L I P R F Y D

GTGACACTTG GAAAAATCTT AGTAGATGGA GTTGATGTAA GAGATTATAA CCTTAAATCA 4320
V T L G K I L V D G V D V R D Y N L K S

CTTCGCCAAA AGATTGGATT TATCCCCCAA AAAGCTCTTT TATTTACAGG GACAATAGGA 4380
L R Q K I G F I P Q K A L L F T G T I G

GAGAATTTAA AATATGGAAA AGCTGATGCT ACTATTGATG ATCTTAGACA AGCGGTTGAT 4440
E N L K Y G K A D A T I D D L R Q A V D

ATTTCTCAAG CTAAAGAGTT TATTGAGAGT CACCAAGAAG CCTTGAAAC GCATTTAGCT 4500
I S Q A K E F I E S H Q E A F E T H L A

GAAGGTGGGA GCAATCTTTC TGGGGGTCAA AAACAACGGT TATCTATTGC TAGGGCTGTT 4560
E G G S N L S G G Q K Q R L S I A R A V

GTTAAAGATC CAGATTTATA TATTTTGTAT GATTCATTTT CTGCTCTCGA TTATAAGACA 4620
V K D P D L Y I F D D S F S A L D Y K T

GACGCTACTT TAAGAGCGCG TCTAAAAGAA GTAACCGGTG ATTCTACAGT TTTGATAGTT 4680
D A T L R A R L K E V T G D S T V L I V

GCTCAAAGGG TGGGTACGAT TATGGATGCT GATCAGATTA TTGTCCTTGA TGAAGGCGAA 4740
A Q R V G T I M D A D Q I I V L D E G E

ATTGTCGGTC GTGGTACCCA CGCTCAATTA ATAGAAAATA ATGCTATTTA TCGTGAAATC 4800
I V G R G T H A Q L I E N N A I Y R E I

GCTGAGTCAC AACTGAAGAA CAAAACTTA TCAGAAGGAG AGTGATTGTA TGAGAAAAAA 4860
A E S Q L K N Q N L S E G E M R K K
|---->

ATCTGTTTTT TTGAGATTAT GGTCTTACCT AACTCGCTAC AAAGCTACTC TTTTCTTAGC	4920
S V F L R L W S Y L T R Y K A T L F L A	
GATTTTTTTG AAAGTTTAT CTAGTTTAT GAGTGTTCTG GAGCCTTTTA TTTTAGGGTT	4980
I F L K V L S S F M S V L E P F I L G L	
AGCGATAACA GAGTTGACTG CTAACCTTGT TGATATGGCT AAGGGAGTTT CTGGGGCAGA	5040
A I T E L T A N L V D M A K G V S G A E	
ATTGAACGTT CCTTATATTG CTGGTATTTT GATTATTTAT TTTTCAGAG GTGTTTCTA	5100
L N V P Y I A G I L I I Y F F R G V F Y	
TGAATTAGGT TCTTATGGCT CAAATT (SEQ ID NO:7)	5126
E L G S Y G S N	

FIG. 2a

NFDIETTTFE	AMKKHASLLE	KISVERSFIE	FDKLLLAPYW	RKGMLALIDS	50
HAFNYLPCLK	NRELQLSAFL	SQLDKDFLFE	TSEQAWASLI	LSMEVEHTKT	100
FLKKWKTSTH	FQKDVEHIVD	VYRIREQMGL	AKEHLYRYGK	TIKQAEGIR	150
KARGLMVDFE	KIEQLDSELA	IHDRHEIVVN	GGTLIKKLG	KPGPQMGDII	200
SQIELAIVLG	QLINEEEAIL	HFVKQYLM	(SEQ ID NO:8)		229

FIG. 2b

MSDFLVDGLT	KSVGDKTVFS	NVSFIIHSLD	RIGIIGVNGT	GKTTLLDVIS	50
GELGFDGDRS	PFSSANDYKI	AYLKQEPDFD	DSQTILDTVL	SSDLREMAI	100
KEYELLLNHY	EESKQSRLEK	VMAEMDSLDA	WSIESEVKTV	LSKLGITDLQ	150
LSVGELSGGL	RRRVQLAQVL	LNDADLLLLD	EPTNHLDIDT	IAWLTNFLKN	200
SKKTVLFITH	DRYFLDNVAT	RIFELDKAQI	TEYQGNQDY	URLRAEQDER	250
DAASLHKKKQ	LYKQELAWMR	TQPQARATKQ	QARINRFQNL	KNDLHQTSDT	300
SDLEMTFETS	RIGKKVINFE	NVSFSYPDKS	ILKDFNLLIQ	NKDRIGIVGD	350
NGVGKSTLLN	LIVQDLQPDS	GNVSIGETIR	VGYFSQQLHN	MDGSKRVINY	400
LQEVADDEVKT	SVGTTSVTEL	LEQFLFPRST	HGTQIAKLSG	GEKKRLLYLLK	450
ILIEKPNVLL	LDEPTNDLDI	ATLTVLENFL	QGFGGPVITV	SHDRYFLDKV	500
ANKIIAFEDN	DIREFFGNYT	DYLDEKAFNE	QNEVISKKE	STKTSREKQS	550
RKRMSYFEKQ	EWATIEDDIM	ILENTITRIE	NDMQTCGSDF	TRLSDLQKEL	600
DAKNEALLEK	YDRYEYLSL	DT	(SEQ ID NO:9)		622

FIG. 2c

MIIRPIIKND	DQAVAQLIRQ	SLRAYDLDP	DTAYSDPHLD	HLTSYYEKIE	50
KSGFFVIEER	DEIIGCGGFG	PLKNLIAEMQ	KVYIAERFRG	KGLATDLVKM	100
IEVEARKIGY	RQLYLETAST	LSRATAVYKH	MGYCALSQPI	ANDQGHTAMD	150
IWMIKDL	(SEQ ID NO:10)				157

FIG. 2d

MAYIWSYLKR	YPNWLWLDLL	GAMLFVTVIL	GMPTALAGMI	DNGVTKGDRT	50
GVYLTWFIMF	IFVVLGIIGR	ITMAYASSRL	TTTMIRDMRN	DMYAKLQEYS	100
HHEYEQIGVS	SLVTRMTSDT	FVLMQFAEMS	LRLGLVTPMV	MIFSVMILI	150
TSPSLAWLVA	VAMPLLGVV	LYVAIKTKPL	SERQQTMLDK	INQYVRENLT	200
GLRVVRA FAR	ENFQSQKFQV	ANQRYTDTST	GLFKLTGLTE	PLFVQIIAM	250
IVAIVWFALD	PLQGAIKIG	DLVAFIEYSF	HALFSFLLFA	NLFTMYPRMV	300
VSSHRIREVM	DMPISINPNA	EGVTDTKLKG	HLEFDNVTFA	YPGETESPVL	350
HDISFKAKPG	ETIAFIGSTG	SGKSSLVNLI	PRFYDVTLGK	ILVDGVDVRD	400
YNLKSLRQKI	GFIPQKALLF	TGTIGENLKY	GKADATIDDL	RQAVDISQAK	450
EFIESHQEAF	ETHLAEGGSN	LSGGQKQRLS	IARAVVKDPD	LYIFDDSFSA	500
LDYKTDATLR	ARLKEVTGDS	TVLIVAQRVG	TIMDADQIIV	LDEGEIVGRG	550
THAQLIENNA	IYREIAESQL	KNQNLSEGE	(SEQ ID NO:11)		579

FIG. 2e

MRKKS VFLRL	WSYLTRYKAT	LFLAIFLKVL	SSFMSVLEPF	ILGLAITELT	50
ANLVDMAKGV	SGAELNVPYI	AGILIIYFFR	GVFYELGSYG	SN	92

(SEQ ID NO:12)

FIG. 2f


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AATTGGAAG TGCTCTATCA ACAGTTGAAG TAAAGGAGAT TATTAGTGAA GAAAACATAT 60
  F G S  A L S  T V E V  K E I  I S E  E N I W
----->
GGTTATATCG GCTCAGTTGC TGCCATTTTA CTAGCTACTC ATATTGGAAG TTACCAACTT 120
  L Y R  L S C  C H F T  S Y S  Y W K  L P T W

GGTAAGCATC ATATGGGTCT AGCAACAAAG GACAATCAGA TTGCCTATAT TGATGACAGC 180
      M G L  A T K  D N Q I  A Y I  D D S
      |----->
AAAGGTAAGG CAAAAGCCCC TAAAACAAAC AAAACGATGG ATCAAATCAG TGCTGAAGAA 240
K G K A  K A P  K T N  K T M D  Q I S  A E E

GGCATCTCTG CTGAACAGAT CGTAGTCAAA ATTACTGACC AAGGCTATGT GACCTCACAC 300
G I S A  E Q I  V V K  I T D Q  G Y V  T S H

GGTGACCATT ATCATTTTTA CAATGGGAAA GTTCCTTATG ATGCGATTAT TAGTGAAGAG 360
G D H Y  H F Y  N G K  V P Y D  A I I  S E E

TTGTTGATGA CGGATCCTAA TTACCGTTTT AAACAATCAG ACGTTATCAA TGAAATCTTA 420
L L M T  D P N  Y R F  K Q S D  V I N  E I L
      |----->
GACGGTTACG TTATTAAAGT CAATGGCAAC TATTATGTTT ACCTCAAGCC AGGTAGTAAG 480
D G Y V  I K V  N G N  Y Y V Y  L K P  G S K

CGCAAAAACA TTCGAACCAA ACAACAAATT GCTGAGCAAG TAGCCAAAGG AACTAAAGAA 540
R K N I  R T K  Q Q I  A E Q V  A K G  T K E

GCTAAAGAAA AAGGTTTAGC TCAAGTGGCC CATCTCAGTA AAGAAGAAGT TCGGCAGTC 600
A K E K  G L A  Q V A  H L S K  E E V  A A V

AATGAAGCAA AAAGACAAGG ACGCTATACT ACAGACGATG GCTATATTTT TAGTCCGACA 660
N E A K  R Q G  R Y T  T D D G  Y I F  S P T

GATATCATTG ATGATTTAGG AGATGCTTAT TTAGTACCTC ATGGTAATCA CTATCATTAT 720
D I I D  D L G  D A Y  L V P H  G N H  Y H Y

ATTCCTAAAA AGGATTTGTC TCCAAGTGAG CTAGCTGCTG CACAAGCCTA CTGGAGTCAA 780
I P K K  D L S  P S E  L A A A  Q A Y  W S Q

AAACAAGGTC GAGGTGCTAG ACCGTCTGAT TACCGCCCGA CACCAGCCCC AGGTCGTAGG 840
K Q G R  G A R  P S D  Y R P T  P A P  G R R

AAAGCCCCAA TTCCTGATGT GACGCCTAAC CCTGGACAAG GTCATCAGCC AGATAACGGT 900
K A P I  P D V  T P N  P G Q G  H Q P  D N G

GGCTATCATC CAGCGCCTCC TAGGCCAAAT GATGCGTCAC AAAACAAACA CCAAAGAGAT 960
G Y H P  A P P  R P N  D A S Q  N K H  Q R D

GAGTTTAAAG GAAAAACCTT TAAGGAACCTT TTAGATCAAC TACACCGTCT TGATTGAAA 1020
E F K G  K T F  K E L  L D Q L  H R L  D L K

TACCGTCATG TGGAAGAAGA TGGGTTGATT TTTGAACCGA CTCAAGTGAT CAAATCAAAC 1080
Y R H V  E E D  G L I  F E P T  Q V I  K S N

GCTTTTGGGT ATGTGGTGCC TCATGGAGAT CATTATCATA TTATCCCAAG AAGTCAGTTA 1140
A F G Y  V V P  H G D  H Y H I  I P R  S Q L

TCACCTCTTG AAATGGAATT AGCAGATCGA TACTTAGCTG GCCAAACTGA GGACAATGAC 1200
S P L E  M E L  A D R  Y L A G  Q T E  D N D

TCAGGTTTCTAG AGCACTCAAA ACCATCAGAT AAAGAAGTGA CACATACCTT TCTTGGTCAT 1260

```

S G S E H S K P S D K E V T H T F L G H
 CGCATCAAAG CTTACGGAAA AGGCTTAGAT GGTAAACCAT ATGATACGAG TGATGCTTAT 1320
 R I K A Y G K G L D G K P Y D T S D A Y
 GTTTTtagta AAGAATCCAT TCATTcagtg GATAAATCAG GAGTTACAGC TAAACACGGA 1380
 V F S K E S I H S V D K S G V T A K H G
 GATCATTTCC ACTATATAGG ATTTGGAGAA CTTGAACAAT ATGAGTTGGA TGAGGTCGCT 1440
 D H F H Y I G F G E L E Q Y E L D E V A
 AACTGGGTGA AAGCAAAAGG TCAAGCTGAT GAGCTTGCTG CTGCTTTGGA TCAGGAACAA 1500
 N W V K A K G Q A D E L A A A L D Q E Q
 GGCAAGAAA AACCCTCTT TGACACTAAA AAAGTGAGTC GCAAAGTAAC AAAAGATGGT 1560
 G K E K P L F D T K K V S R K V T K D G
 AAAGTGGGCT ATATGATGCC AAAAGATGGT AAGGACTATT TCTATGCTCG TGATCAACTT 1620
 K V G Y M M P K D G K D Y F Y A R D Q L
 GATTGACTC AGATTGCCTT TGCCGAACAA GAACTAATGC TTAAGATAA GAAGCATTAC 1680
 D L T Q I A F A E Q E L M L K D K K H Y
 CGTTATGACA TTGTTGACAC AGGTATTGAG CCACGACTTG CTGTAGATGT GTCAAGTCTG 1740
 R Y D I V D T G I E P R L A V D V S S L
 CCGATGCATG CTGGTAATGC TACTTACGAT ACTGGAAGTT CGTTTGTTAT CCCACATATT 1800
 P M H A G N A T Y D T G S S F V I P H I
 GATCATATCC ATGTCGTTCC GTATTcattg TTGACGCGCG ATCAGATTGC AACAGTCAAG 1860
 D H I H V V P Y S W L T R D Q I A T V K
 TATGTGATGC AACACCCCGA AGTTCGTCCT GATGTATGGT CTAAGCCAGG GCATGAAGAG 1920
 Y V M Q H P E V R P D V W S K P G H E E
 TCAGGTTCCG TCATTCCAAA TGTTACGCCT CTTGATAAAC GTGCTGGTAT GCCAAACTGG 1980
 S G S V I P N V T P L D K R A G M P N W
 CAAATTATCC ATTCTGCTGA AGAAGTTCAA AAAGCCCTAG CAGAAGGTCG TTTTGCAACA 2040
 Q I I H S A E E V Q K A L A E G R F A T
 CCAGACGGCT ATATTTTCGA TCCACGAGAT GTTTTGCCCA AAGAACTTT TGTATGAAAA 2100
 P D G Y I F D P R D V L A K E T F V W K
 GATGGCTCCT TTAGCATCCC AAGAGCAGAT GGCAGTTCAT TGAGAACCAT TAATAAATCT 2160
 D G S F S I P R A D G S S L R T I N K S
 GATCTATCCC AAGCTGAGTG GCAACAAGCT CAAGAGTTAT TGGCAAAGAA AAATACTGGT 2220
 D L S Q A E W Q Q A Q E L L A K K N T G
 GATGCTACTG ATACGGATAA ACCCAAAGAA AAGCAACAGG CAGATAAGAG CAATGAAAAC 2280
 D A T D T D K P K E K Q Q A D K S N E N
 CAACAGCCAA GTGAAGCCAG TAAAGAAGAA AAAGAATCAG ATGACTTTAT AGACAGTTTA 2340
 Q Q P S E A S K E E K E S D D F I D S L
 CCAGACTATG GTCTAGATAG AGCAACCCTA GAAGATCATA TCAATCAATT AGCACAACAAA 2400
 P D Y G L D R A T L E D H I N Q L A Q K
 GCTAATATCG ATCCTAAGTA TCTCATTTTC CAACCAGAAG GTGTCCAATT TTATAATAAA 2460
 A N I D P K Y L I F Q P E G V Q F Y N K

AATGGTGAAT TGGTAACTTA TGATATCAAG ACACTTCAAC AAATAAACCC TTAACCAAAA 2520
 N G E L V T Y D I K T L Q Q I N P
 GAAGATCTCA TTGTTAAAGC ACTGCTTTGT CAAAGCAAGT TACGGTGATT TTGAAGTCAT 2580
 TCTATGTAAC GAGTAGTGAT AAAAGTTGGA TAATAGCGGT TTTCTTTTGC AAAGAAATGG 2640
 TATCCATGTT AGAATAGTAA AAAAAGAGGA GGATTCTTGG ACTAATGTCA AATAAGTAGA 2700
 CAGAAAACCTG TGTTATTTTA TTGCGTTAAA ATAATTTTCT TCTTTCTGAT TAGGGGTTAG 2760
 .K I A N F Y N E E K Q N P T L
 TCCTAGATTA GCCGTATGTG GGTTGTAATT GTTATAAAAA TTCTCAATGT ATTCAAAGCA 2820
 G L N A T H P N Y N N Y F N E I Y E F C
 GTCTAATTGA ACCTGTTTGA TATTTTGATA ATGTTTTCGG TTGATTGTGC TATGCTTTAA 2880
 D L Q V Q K I N Q Y H K R N I Q R H K L
 ATACTTGAAG AATGCTTCAG TTACGGCATT ATCATAAGGA TATCCAGGAT TAGAAAAAGA 2940
 Y K F F A E T V A N D Y P Y G P N S F S
 ATGCATGATA TTGGCACTGC ACCCTAATAG TGAGACGCAA GAAAAACACT TTTAGGCAAT 3000
 H M A I
 <----|
 CAGTTTTCTG TACTGTACAG GCGACTGGTC GTTAAATCTC TGTTGAATTC TAGTTTCATT 3060
 L K R Y Q V P S Q D N L R Q Q I R T E N
 ATAAATGTA ATGTAATTTT TAACAATATT TGTTATACTA TCTTTGTGTG ATTTTCTCCT 3120
 Y F T I Y N K V I N T I S D K N Y K R R
 ATTATGGAAA TAAAAGGTTT CAGTCTTTAG GACGGTGTGA AACCATTCAA TACAGGCATT 3180
 N H F Y F T E T K L V T H F W E I C A N
 ATCTGCAGGT GTTCCTTTTC GAGACATTGA GCGGATAATG TCTTTTCCG TGCAAGCCTG 3240
 D A P T G K R S M S R I I D K E T C A Q
 GTAGTAAGCC ATAGAAGTAT ACACTGAGCC TTGGTCACTG TGTAAGATTG CTCCTTTATT 3300
 Y Y A M
 <----|
 TAGGCAATTT TAACGTATTA AGGGTGTCTA GTACAAAATC CGTGTCTGTA CAATCTGAGA 3360
 K P L K L Q N L T D L V F D T D Q C D S
 TAGTGTAGC TATAATTTCT CGGTTATAGA GATTGATAAT TGATGAGAGA TACAATTTAC 3420
 I T Y A I I E R N Y L N M I S S L Y L K
 AGTTACCGAA ATATAGGTAG GTAATATCTG TTACGAGCTT TTCCTTAGGC TTATCGGCAT 3480
 C N G F Y L Y T I D T V L K E K P K D A
 GGAAATCCCG ACTCAATTTA TTATCTGTTA AATAATAAGC TTTACCCAAA TTGGGAACCT 3540
 H G D R S L K N D T L Y Y A K G L N P V
 TCTTGGTACG TGTCCGACAA AGCCAGCCAT TATTTTTCAT GATACGATAG ACTTTCTTTG 3600
 K K T R T R C L W G N N K M I R Y V K K
 TATTAACAGT CAATCCGTGG ATTTTGTGA GCAATCGTGT AATGGTACGA TAGCCATAAA 3660
 T N V T L G H I K K L L R T I T R Y G Y
 TAAAGTGATT CTCCATACAG AGCTGTTCAA TTAATTCAAT AAGGTCATCT TTTTTCGCG 3720
 I F H N E M
 <----|

CTTCATAC TCCTTTTCC AACGGTAATA GGTCGACCGC TTGACCTTAA AACAGTCTAG 3780
 AATGAAACT ATCGGGTAGT TGTTTTTATA GTCTTCCACA AGCTTGATAA GACTTACTTT 3840
 ATCGATTTC TTATCAAGCC TCGATACTTT TTTAAGAGGT CAACCTGTAA TTGTAATTGT 3900
 I S K R I L G R Y K K L L D V Q L Q L Q
 TCCACTTCAG ACAGATGTTT CAAGCCTTTA CCGTAGGTAT ATTGCTTGCC AACACCTTGA 3960
 E V E S L H E L G K G Y T Y Q K G V G Q
 TGAAAACGAT AAAGCTCCTC GTTTTCGTAC CATTTCATCC AAGTATAGAT TTGACTATTA 4020
 H F R Y L E E N E Y W K M W T Y I Q S N
 TTTTGTATGC CTAAAGTCTC CATAATAACT CTGTAGACT TGCCTGCTTT CTTCATATCG 4080
 N K I G L T E M I V R N S K G A K K M D
 ATGCAAGCCA GCTTAGTTTC CCATGAATAT GCTTTTTTAA CCATAATAAA ACATTCCTGT 4140
 I C A L K T E W S Y A K K V M
 TTCTAGTTTA CTAAATTTC ACAGGAGTGT TTTCTTTTG TCTCATTTTA GGGATTCACT 4200
 GCCTATTGTT GTCATCAATT ATTTTCTAA ATTCCCGGA CTAAATTGT GACCCTTGGT 4260
 CGGAATGAAA GAGAAGTGT CTTCAATCT TTCTTTTATT AAGTGAAAAG GCAACACTTT 4320
 TCTGTACAAC ATTTATAAAG TGTTTTTCTA GGCAATTAAT CTTTGTAGTCA TTGGTGTTTG 4380
 . A I L R K T M P T Q
 GTAGTTGAGA CTACCATGAA TGCGGTGGTA ATTCCACCAA TGAACATAGT CTTTAGTCTT 4440
 Y N L S G H I R H Y N W W H V Y D K T K
 AAGAGCTAGT TCTTCCAGCA ATTGAAAGGT TTCTTGATAA ACAAATTCAA TTTTGAAGC 4500
 L A L E E L L Q F T E Q Y V F E I K F A
 ACGATACGTA CTTTCAGCTA CGGCATTGTC ATAAGGATAA CCAGCCTGAC TAAGCGAACG 4560
 R Y T S E A V A N D Y P Y G A Q S L S R
 TGTGATTCCA AAGGCTTCCA ATATTTCATC AATTAAGTGA TTATCAAACT CTTTGCCACG 4620
 T I G F A E L I E D I L Q N D F E K G R
 ATCTGAATGG AACATCTTGA CTTTGGTCAG GGCGTAAGGG ATGCTTTGTA TGGCTTGCTT 4680
 D S H F M K V K T L A Y P I S Q I A Q K
 AACGAGTTCA GCGGTCTTGT GCCAACCAAG AGACAGGCCG ATGATTTCAC GGTGTATAG 4740
 V L E A T K H W G L S L G I I E R N Y L
 GTCAATGATG AGGCAAACAT AAGCCCAACG ATTGCCTACA CGAACATAGG TTAAGTCAGT 4800
 D I I L C V Y A W R N G V R V Y T L D T
 GACTAAGGCT TGTAGTGGTC TTTCTTGCTT AAATTGCCTG TCTAAGTGGT TGGGAATAGG 4860
 V L A Q L P R E Q K F Q R D L H N P I P
 GGCTTCATTC TTGCCTCTAG AATGTGGTTT GAAGGTGGCT TTCTGATAAA CAGAAACCAA 4920
 A E N K G R S H P K F T A K Q Y V S V L
 ATTGAGTCGC TTCATAATGC GTCGAATCCG ACGACGTGAA AGTGTGATAC CTTGTTTATT 4980
 N L R K M I R R I R R R S L T I G E N N
 CAAGCATATT TTGATTTTTC TGGATCCGTA TCTAGACTCG CTATCGAGAA AAATTCCTTT 5040
 L C I K I K R S G Y R S E S D L F I R K

AATAGTTTCT TCAAACCTCCG TTTCAGATAC TGACTCCACG GCTTGATAGT AATAACTTGA 5100
 I T E E F E T E S V S E V A Q Y Y Y S S
 GTGTGGCATA TTCAGCCAGC GACACATCTT TGAAATGCTG TATTTATCCT TATTAGCAGT 5160
 H P M N L W R C M K S I S Y K D K N A T
 GATTATTTCC CTTTTGTGC CATAATCACC GCTGCTTGCT TTAGGATATC TAATT 5215
 I I E R K T G Y D G S S A K P Y R I
 (SEQ ID NO:13) <-----|

FIG. 3a

FGSALSTVEV KEIISEENIW LYRLSCCHFT SYSYWKLPW 40
 (SEQ ID NO:14)

FIG. 3b

MGLATKDNQI AYIDDSKGKA KAPKTNKTMD QISAEEGISA EQIVVKITDQ 50
 GYVTSBGDHY HFYNGKVPYD AIISEELLMT DPNYRFKQSD VINEILDGYV 100
 IKVNGNYVVY LKPGSKRKNI RTKQQIAEQV AKGTKEAKEK GLAQVAHLSK 150
 EEVAAVNEAK RQGRYTTDDG YIFSPTDIID DLGDAYLVPH GNHYHYIPKK 200
 DLSPSELAAA QAYWSQKQGR GARPSDYRPT PAPGRRKAPI PDVTPNPGQG 250
 HQPDNGGYHP APPRPNDASQ NKHQDEFKQ KTFKELLDQL HRLDLKYRHV 300
 EEDGLIFEPT QVIKSNAFGY VVPHGDHYHI IPRSQLSPLE MELADRYLAG 350
 QTEDNDGSGSE HSKPSDKEVT HTFLGHRIKA YGKGLDGKPY DTSDAYVFSK 400
 ESIHSDVKSG VTAKHGDHFI YIGFGELEQY ELDEVANWVK AKGQADELAA 450
 ALDQEQGKEK PLEDTKKVSF KVTGDKGVGY MMPKDGKDYF YARDQLDLTQ 500
 IAFAEQELML KDKKHRYDI VDTGIEPRLA VDVSSLPMHA GNATYDTGSS 550
 FVIPHIDHIH VVPYSWLTRD QIATVKYVMQ HPEVREPDVWS KPGHEESGSV 600
 IPNVTPLDKR AGMPNWQIIH SAEVQKALA EGRFATPDGY IFDPRDLAK 650
 ETFVWKDGSF SIPRADGSSL RTINKSDLSQ AEWQQAQELL AKKNTGDATD 700
 TDKPKEKQQA DKSNNQQPS EASKEEKESD DFIDSLPDYG LDRATLEDHI 750
 NQLAQKANID PKYLIFQPEG VQFYNGKNGEL VTYDIKTLQQ INP 793
 (SEQ ID NO:15)

FIG. 3c

MTDPNYRFKQ	SDVINEILDG	YVIKVNNGYY	VYLKPGSKRK	NIRTKQQIAE	50
QVAKGTKEAK	EKGLAQVAHL	SKEEVAAVNE	AKRQGRYTTD	DGYIFSPTDI	100
IDDLGDAYLV	PHGNHYHYIP	KKDLSPSELA	AAQAYWSQKQ	GRGARPSDYR	150
PTPAPGRRKA	PIPDVTPNPG	QGHQPDNGGY	HPAPPRPND	SONKHQRDEF	200
KGKTFKELLD	QLHRLDLKYR	HVEEDGLIFE	PTQVIKSNF	GYVVPBGDHY	250
HIIPRSQSLP	LEMELADRYL	AGQTEDNDG	SEHSKPSDKE	VTHTFLGHRI	300
KAYGKGLDGK	PYDTSDAYVF	SKESIHSVDK	SGVTAKHGDH	FHYIGFGELE	350
QYELDEVANW	VKAKGQADEL	AAALDQEQGK	EKPLFDTKKV	SRKVTKDGKV	400
GYMMPKDGKD	YFYARDQLDL	TQIAFAEQEL	MLKDKKHRY	DIVDTGIEPR	450
LAVDVSSLPM	HAGNATYDTG	SSFVIPHIDH	IHVVPYSWLT	RDQIATVKYV	500
MQHPEVRPDV	WSKPGHEESG	SVIPNVTPLD	KRAGMPNWQI	IHSAAEVQKA	550
LAEGRFATPD	GYIFDPRDVL	AKETFVWKDG	SFSIPRADGS	SLRTINKSDL	600
SQAEWQQAQE	LLAKKNTGDA	TDTDKPKEKQ	QADKSNENQQ	PSEASKEEKE	650
SDDFIDSLPD	YGLDRATLED	HINQLAQKAN	IDPKYLIFQP	EGVQFYNNKG	700
ELVTYDIKTL	QQINP	(SEQ ID NO:16)			715

FIG. 3d

MHSFSNPGYP	YDNAVTEAFF	KYLKHRQINR	KHYQNIQVQ	LDCFYIENF	50
YNNYNPHTAN	LGLTPNQKEE	NYFNAIK	(SEQ ID NO:17)		77

FIG. 3e

MAYYQACTEK	DIIRMSRKG	TPADNACIEW	FHTVLKTETF	YFHNRRKYNK	50
DSITNIVKNY	ITFYNETRIQ	QRLNDQSPVQ	YRKLIA	(SEQ ID NO:18)	86

FIG. 3f

MENHFIYGYR	TITRLLKKIH	GLTVNTKKVY	RIMKNNGWLC	RTRTKKVPNL	50
GKAYYLTDNK	LSRDFHADKP	KEKLVDITY	LYFGNCKLYL	SSIMNLYNRE	100
IIAYTISDCQ	DTDFVLDTLN	QLKLPK	(SEQ ID NO:19)		126

FIG. 3g

MVKKAYSWET KLACIDMKKA GKSNRVIMET LGIKNNSQIY TWMKWYENEE 50
 LYRFHQGVGK QYTYGKGLEH LSEVEQLQLQ VDLLKKYRGL IRKSIK 96
 (SEQ ID NO:20)

FIG. 3h

IRYPKASSGD YGTKREIITA NKDKYSISKM CRWLNMPHSS YYYQAVESVS 50
 ETEFEETIKR IFLDSESRYG SRKIKICLNN EGITLSRRRI RRIMKRLNLV 100
 SVYQKATFKP HSRGKNEAPI PNHLDRQFKQ ERPLQALVTD LTYVRVGNRW 150
 AYVCLIIDLY NREIIGLSLG WHKTAELVKQ AIQSIPYALT KVKMFHSDRG 200
 KEFDNQLIDE ILEAFGITRS LSQAGYPYDN AVAESTYRAF KIEFVYQETF 250
 QLLEELALKT KDYVHWWNYH RIHGSLNYQT PMTKRLIA (SEQ ID NO:21)288

FIG. 3i

AATTTGAAAG CAGAATTATC TGTAAGAGAT GAGCAATATA CAGCAACAGT TTATGGTAAA 60
 N L K A E L S V E D E Q Y T A T V Y G K
 ---->
 TCTGCTCATG GTTCAACACC ACAAGAAGGT GTTAATGGGG CGACTTATTT AGCTCTTTAT 120
 S A H G S T P Q E G V N G A T Y L A L Y
 CTAAGTCAAT TTGATTTTGA AGGTCCTGCT CGTGCTTTCT TAGATGTTAC AGCCAACATT 180
 L S Q F D F E G P A R A F L D V T A N I
 ATTCACGAAG ACTTCTCAGG TGAAAACTT GGAGTAGCTT ATGAAGATGA CTGTATGGGA 240
 I H E D F S G E K L G V A Y E D D C M G
 CCATTGAGCA TGAATGCAGG TGTCTCCAG TTTGATGAAA CTAATGATGA TAATACTATC 300
 P L S M N A G V F Q F D E T N D D N T I
 GCTCTTAATT TCCGTTACCC ACAAGGGACA GATGCTAAAA CTATCCAAAC TAAGCTTGAG 360
 A L N F R Y P Q G T D A K T I Q T .K L E
 AAACCTTAACG GAGTTGAAAA AGTGACTCTT TCTGACCATG AACACACACC AACTATGTA 420
 K L N G V E K V T L S D H E H T P H Y V
 CCTATGGACG ATGAATTAGT ATCAACCTTA CTAGCTGTCT ATGAAAAGCA AACTGGTCTT 480
 P M D D E L V S T L L A V Y E K Q T G L
 AAAGGACATG AACAGGTTAT TGGTGGTGGG ACATTTGGTC GCTTACTTGA ACGGGGTGTT 540
 K G H E Q V I G G G T F G R L L E R G V
 GCATACGGTG CCATGTTCCC AGGAGATGAA AACACTATGC ATCAAGCTAA TGAGTACATG 600
 A Y G A M F P G D E N T M H Q A N E Y M
 CCTTTAGAAA ATATTTTCCG TTCGGCTGCT ATCTACGCAG AAGCTATCTA TGAATTAATC 660

P L E N I F R S A A I Y A E A I Y E L I
 AAATAAAATA ATCCTTAAAC TAAATATGTG ATCAATGATA AAGGGTGGTG AAGACATGAA 720
 K .
 AGTGTCTTTG CCTCTTTTCA TAAGGTTAGA TTTGGAGACT TTATGACTGA CTTGGAAAAA 780
 M T D L E K
 |---->
 ATTATTAAAG CAATAAAAAG TGATTCACAG AATCAAAATT ATACAGAAAA TGGTATTGAT 840
 I I K A I K S D S Q N Q N Y T E N G I D
 CCTTGTGTTG CTGCTCCTAA AACAGCTAGG ATCAATATTG TTGGCCAAGC ACCTGGTTTA 900
 P L F A A P K T A R I N I V G Q A P G L
 AAAACTCAAG AAGCAAGACT CTATTGGAAA GATAAATCTG GAGATCGTCT ACGCCAGTGG 960
 K T Q E A R L Y W K D K S G D R L R Q W
 CTTGGAGTTG ATGAAGAGAC ATTTTACCAT TCTGGAAAAT TTGCTGTTTT ACCTTTAGAT 1020
 L G V D E E T F Y H S G K F A V L P L D
 TTTTATTACC CAGGCAAAGG AAAATCAGGA GATTTACCCC CTAGAAAAGG TTTTGGCGAG 1080
 F Y Y P G K G K S G D L P P R K G F A E
 AAATGGCACC CTCTTATTTT AAAAGAAATG CCTAATGTTC AATTGACCTT GCTAGTTGGT 1140
 K W H P L I L K E M P N V Q L T L L V G
 CAGTATGCTC AGAAATATTA TCTTGAAGC TCCGCACATA AAAATCTAAC AGAAACAGTT 1200
 Q Y A Q K Y Y L G S S A H K N L T E T V
 AAAGCTTACA AAGACTATCT ACCCGATTAT TTACCCCTGG TTCACCCATC ACCGCGAAAT 1260
 K A Y K D Y L P D Y L P L V H P S P R N
 CAAATTTGGC TAAAGAAGAA TCCATGGTTT GAAAAAGATC TAATCGTTGA TTTACAAAAG 1320
 Q I W L K K N P W F E K D L I V D L Q K
 ATAGTAGCAG ATATTTTAAA AGATTAAGGA TAGGAGTTGG TATGAGAGAT AATCATCTAC 1380
 I V A D I L K D .
 M R D N H L H
 |---->
 ACACGTATTT TTCCTATGAT TGTCAAACGG CATTGAGGA CTATATTAAT GGTTTACAG 1440
 T Y F S Y D C Q T A F E D Y I N G F T G
 GTGAATTTAT CACGACAGAA CATTTTGATT TATCAAATCC TTACACCGGT CAAGACGATG 1500
 E F I T T E H F D L S N P Y T G Q D D V
 TTCCTGATTA TAGTGCTTAT TGTCAAAAAA TAGATTATCT TAATCAGAAA TATGGAAATC 1560
 P D Y S A Y C Q K I D Y L N Q K Y G N R
 GATTTAAAAA AGGAATTGAA ATCGGTTATT TTAAAGATAG GGAATCAGAT ATTTTAGATT 1620
 F K K G I E I G Y F K D R E S D I L D Y
 ATTTAAAAA TAAAGAATTT GATTTAAAC TATTGTCAAT CCATCATAAT GGTAGGTATG 1680
 L K N K E F D L K L L S I H H N G R Y D
 ATTATCTGCA AGAAGAAGCT CTGAAAGTAC CAACAAAGGG AGCTTTTAGC AGATTACTTT 1740
 Y L Q E E A L K V P T K G A F S R L L .
 AATCGTATGG AATTTGCCAT AGGCCGTGTG GAAGCGCAGC TTTTAGCTCA CTTTGATTAT 1800
 GGTTTTCGTA AGTTAACTT AGATGTAGAA GATTTAAAC CGTTTGAAAC GCAATTGAAG 1860
 CGCATTTTCA TAAAGATGTT ATCTAAGGGG TTAGCTTTTG AACTAAATAC CAAATCCCTT 1920

22/40

AAACTAACG CCAAAGAGAT TGGTTTTCTA ATCTTAGATG AAAGTAAGAC AGGAGATGCA 3300
 K T N A K E I G F L I L D E S K T G D A
 GTGAAAGTTC AACCCAACGA CTATGTTTTT AGAGATTTAG CTAACCATAA CCAAATTTTT 3360
 V K V Q P N D Y V F R D L A N H N Q I F
 GTAAAGATA AGGATCCAAA GGGTTATAAT AATCCTTATT ACATTGATCA AGTGCAGCTA 3420
 V K D K D P K V Y N N P Y Y I D Q V Q L
 AAGGATGCCC AACAAATTGA TTTAACAAGT ATTCAAGCAA GTTTTACAAC TCTAGATGGG 3480
 K D A Q Q I D L T S I Q A S F T T L D G
 GTAGATAAAA CTGAAATTTT AAAAGAATTG AAAGTGACTG ATAAAAATCA AAATGCTATA 3540
 V D K T E I L K E L K V T D K N Q N A I
 CAAATTTCTG ATATCACTCT CGATACTAGT AAATCTCTTT TAATAATCAA AGGCGACTTT 3600
 Q I S D I T L D T S K S L L I I K G D F
 AATCCTAAAC AAGGTCATTT CAACATATCT TATAATGGTA ACAATGTCAT GACAAGGCAA 3660
 N P K Q G H F N I S Y N G N N V M T R Q
 TCTTGGGAAT TTAAAGACCA ACTTTATGCT TATAGTGGAA ATTTAGGTGC AGTTCTCAAT 3720
 S W E F K D Q L Y A Y S G N L G A V L N
 CAAGATGGTT CAAAAGTTGA AGCCAGCCTC TGGTCACCGA GTGCTGATAG TGTCACTATG 3780
 Q D G S K V E A S L W S P S A D S V T M
 ATTATTTATG ACAAAGATAA CCAAAACAGG GTTGTAGCGA CTACCCCCCT TGTGAAAAAT 3840
 I I Y D K D N Q N R V V A T T P L V K N
 AATAAAGGTG TTTGGCAGAC GATACTTGAT ACTAAATTAG GTATTAAAAA CTATACTGGT 3900
 N K G V W Q T I L D T K L G I K N Y T G
 TACTATTATC TTTACGAAAT AAAAAGAGGT AAGGATAAGG TTAAGATTTT AGATCCTTAT 3960
 Y Y Y L Y E I K R G K D K V K I L D P Y
 GCAAAGTCAT TAGCAGAGTG GGATAGTAAT ACTGTTAATG ATGATATTAA AACGGCTAAA 4020
 A K S L A E W D S N T V N D D I K T A K
 GCAGCTTTTG TAAATCCAAG TCAACTTGGG CCTCAAAATT TAAGTTTTCG TAAAATTGCT 4080
 A A F V N P S Q L G P Q N L S F A K I A
 AATTTTAAAG GAAGACAAGA TGCTGTTATA TACGAAGCAC ATGTAAGAGA CTTCACTTCT 4140
 N F K G R Q D A V I Y E A H V R D F T S
 GATCGATCTT TGGATGGAAA ATTAAAAAAT CAATTTGGTA CCTTTGCAGC CTTTTCAGAG 4200
 D R S L D G K L K N Q F G T F A A F S E
 AAAGTAGATT ATTTACAGAA ATTAGGAGTT ACACACATTC AGCTTTTACC GGTATTGAGT 4260
 K L D Y L Q K L G V T H I Q L L P V L S
 TATTTTTATG TTAATGAAAT GGATAAGTCA CGCTCAACAG CTTACACTTC CTCAGACAAT 4320
 Y F Y V N E M D K S R S T A Y T S S D N
 AATTACAATT GGGGCTATGA CCCACAGAGC TATTTTGCTC TTTCTGGGAT GTATTCAGAG 4380
 N Y N W G Y D P Q S Y F A L S G M Y S E
 AAACCAAAAG ATCCATCAGC ACGTATCGCC GAATTAAAC AATTAATACA TGATATTCAT 4440
 K P K D P S A R I A E L K Q L I H D I H

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AAACGTGGCA TGGGGGTTAT ACTTGATGTC GTCTATAATC AACTGCAAA AACTTATCTC 4500
K R G M G V I L D V V Y N H T A K T Y L

TTTGAGGATA TAGAACCTAA TTATTATCAC TTTATGAATG AAGATGGTTC ACCAAGAGAA 4560
F E D I E P N Y Y H F M N E D G S P R E

AGTTTTGGAG GGGGACGTTT AGGAACCACT CATGCAATGA GTCGTCGTGT TTTGGTTGAT 4620
S F G G G R L G T T H A M S R R V L V D

TCCATTAAAT ATCTTACAAG TGAATTTAAA GTTGATGGTT TCCGTTTGA TATGATGGGA 4680
S I K Y L T S E F K V D G F R F D M M G

GATCATGATG CGGCTGCGAT TGAATTAGCT TATAAAGAAG CTAAAGCTAT TAATCCTAAT 4740
D H D A A A I E L A Y K E A K A I N P N

ATGATTATGA TTGGTGAGGG CTGGAGAACA TTCCAAGGCG ATCAAGGTCA GCCGGTTAAA 4800
M I M I G E G W R T F Q G D Q G Q P V K

CCAGCTGACC AAGATTGGAT GAAGTCAACC GATACAGTTG GCGTCTTTTC AGATGATATT 4860
P A D Q D W M K S T D T V G V F S D D I

CGTAATAGCT TGAATCTGG TTTTCCAAAT GAAGGTACTC CAGCTTTCAT CACAGGTGGC 4920
R N S L K S G F P N E G T P A F I T G G

CCACAATCTT TACAAGGTAT TTTTAAAAAT ATCAAAGCAC AACCTGGGAA TTTTGAAGCA 4980
P Q S L Q G I F K N I K A Q P G N F E A

GATTCGCCAG GAGATGTGGT GCAGTATATT GCTGCACATG ATAACCTTAC CTGTCATGAT 5040
D S P G D V V Q Y I A A H D N L T L H D

GTGATTGCAA AATCAATT (SEQ ID NO:22) 5058
V I A K S I .

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FIG. 4a

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NLKAELSVED EQYTATVY GK SAHGSTPQEG VNGATYLALY LSQDFDEGPA 50
RAFLDVTANI IHEDFSGEKL GVAYEDDCMG PLSMNAGVFQ FDETNDNDNTI 100
ALNFRYPQGT DAKTIQTKLE KLNGVEKVTL SDHEHTPHYV PMDDELVSTL 150
LAVYEKQTGL KGHEQVIGGG TFGRLRLRGV AYGAMFPGDE NTMHQANEYM 200
PLENIFRSAA IYAEAIYELI K (SEQ ID NO:23) 221

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FIG. 4b

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MTDLEKIIKA IKSDSQNQNY TENGIDPLFA APKTARINIV GQAPGLKTQE 50
ARLYWKDKSG DRLRQWLGV D EETFYHSGKF AVLPLDFYYP GKGKSGDLPP 100
RKGFAEKWHP LILKEMPNVQ LTLLVGQY AQ KYYLGS SAHK NLTETVKAYK 150
DYLDPYLPV HPSPRNQIWL KKNPWFEKDL IVDLQKIVAD ILKD 194
(SEQ ID NO:24)

```

FIG. 4c

MRDNHLHTYF SYDCQTAFED YINGFTGEFI TTEHFDLSNP YTGQDDVPDY	50
SAYCQKIDYL NQKYGNRFKK GIEIGYFKDR ESDILDYLN KEFDLKLLSI	100
HHNGRYDYLQ EEALKVPTKG AFSRL (SEQ ID NO:25)	126

FIG. 4d

MKRKDLFGDK QTQYTIRKLS VGVASVTTGV CIFLHSPQVF AEEVSVSPAT	50
TAIAESNINQ VDNQQSTNLK DDINSNSETV VTPSDMPDTK QLVSEDTDTQ	100
KGVTEPDKAT SLLEENKGPV SDKNTLDLKV APSTLQNTPD KTSQAIGAPS	150
PTLKVANQAP RIENGYFRLH LKELPQGHPV ESTGLWIWGD VDQPSSNWP	200
GAIPMTDAKK DDYGYVDFK LSEKQRKQIS FLINNAGTN LSGDHHIPLL	250
RPEMNQVWID EKYGIHTYQP LKEGYVRINY LSSSSNYDHL SAWLEKDVAT	300
PSTTWPDGSN FVNQGLYGRY IDVSLKTNK EIGFLILDES KTGDAVKVQP	350
NDYVFRDLAN HNQIFVKDKD PKVYNNPYYI DQVQLKDAQQ IDLTSIQASF	400
TTLDGVDKTE ILKELKVTDK NQNAIQISDI TLDTSKSLLI IKGDFNPKQG	450
HFNISYNGNN VMTRQSWEFK DQLYAYSGNL GAVLNQDGSK VEASLWSPSA	500
DSVTMIYDK DNQNRVATT PLVKNNKGVW QTILDTKLG I KNYTGYYYLY	550
EIKRGKDKVK ILDPYAKSLA EWDSNTVNDD IKTAKAFFVN PSQLGPNLS	600
FAKIANFKGR QDAVIYEAHV RDFTSDRSLD GKLKNQFGTF AAFSEKLDYL	650
QKLGVTHIQL LPVLSYFYVN EMDKSRSTAY TSSDNNYNWG YDPQSYFALS	700
GMYSEKPKDP SARIAELKQL IHDHKGGMG VILDVVYNHT AKTYLFEDIE	750
PNYYHFMNED GSPRESFGGG RLGTTHAMSR RVLVDSIKYL TSEFKVDGFR	800
FDMMGDHDA AIELAYKEAK AINPNMIMIG EGWRTFQGDQ GQPVKPADQD	850
WMKSTDTVGV FSDDIRNSLK SGFPNEGTPA FITGGPQSLQ GIFKNIKAQP	900
GNFEADSPGD VVQYIAAHDN LTLHDVIAKS I (SEQ ID NO:26)	931

FIG. 4e

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AATTCAAAGT TTGACAGAAG GTCAACTTCG TTCTGATATC CCTGAGTTCC GTGCTGGTGA 60
I Q S L T E G Q L R S D I P E F R A G D
----->
TACTGTACGT GTTCACGCTA AAGTTGTTGA AGGTACTCGC GAACGTATTC AGATCTTTGA 120
T V R V H A K V V E G T R E R I Q I F E
AGGTGTTGTT ATCTCACGTA AAGGTCAAGG AATCTCAGAA ATGTACACAG TACGTAAAAT 180
G V V I S R K G Q G I S E M Y T V R K I
TTCTGGTGGT ATCGGTGTAG AGCGTACATT CCCAATTCAC ACTCCTCGTG TTGATAAAAT 240
S G G I G V E R T F P I H T P R V D K I
CGAAGTTGTT CGTTATGGTA AAGTACGTCG TGCTAAACTT TACTACTTAC GCGCATTGCA 300
E V V R Y G K V R R A K L Y Y L R A L Q
AGGTAAAGCT GCACGTATTA AAGAAATCCG TCGTTAATTT TGATGATCAG ATTTTAAAAA 360
TGCTTGGTTG TTTGAGGATA GTAACATATGT TTTAAACTG GACAACCAAG ACGTAAAAAA 420
TCTGCCTGTG GGCAGTTTTT TTACTAGGTC CCCTTAGTTC AATGGATATA ACAACTCCCT 480
. H I Y C S G
CCTAAGGAGT AATTGCTGGT TCGATTCCGG CAGGGGACAT ATTCATTGCA TGTAATAGC 540
G L S Y N S T R N R C P V Y E N C T F L
GGTTAGAGC TATTTTGCCC CAAATTCTC TGATTAAGTT TATCGTTCCT ATCTTTTGT 600
P K S S N Q G L N R Q N L K D N R D K Q
TCTTGTAATT GATGTGCGTA AACTTCTAAA GTGATATTTA AATTCTCGTG ATCTAAACT 660
E Q L Q H A Y V E L T I N L N E H D L V
TGAGAGATGG AAATTAGATA GCTTGCAAAT GTATGCCTGA GAGAGTGCAC TCGTACCTCG 720
Q S I S I L Y S A F T H R L S H V R V E
CGACCAGTTA TTTTTCGGAT AGTTTATTG ACTGCATTAT TTGAAAGTTT GTCGAATAAT 780
R G T I K R I T K N V A N N S L K D F L
CTGTCGTTTT TATTTTTTGT AAATTCATGC AAAAAAATA ATGTATCATT GTCAATTGGT 840
R D N K N K T F E H L F F L T D N D I P
ATATTTCTGA TACTACTTTT GTTTTTTGT GGCAGGTATC TTTGGTTGAA ATGATAATCC 900
I N R I S S K N K T P L Y R Q N F H Y D
CAAGTTTTAT TAATTGATAA ATATTTGTTA GTGTAATCAA TATCATTAAC TGTTAAACCT 960
W T K N I S L Y K N T Y D I D N V T L G
AAACATTCAG CGAAGCGCAT GCCAGTTTTA GCGATGAGGT ATAACGCTGC ATACGATTGA 1020
L C E A F R M
<-----|
TGTTGTGATT TTTCTTTACA AATTTTTATC AAGCGTAAGT ATTCATTGGT TTCAAGAAAT 1080
TTTATCTCTA TTTACGCCCC TTATTTTTTG CTTTAACCTT AGTGAATAAA CAAAAATTTT 1140
TTTCTATATA TCCCTCGTGA ACAGCCATGG ATACGCAGGC TTTTACATGT ATGTTAAAC 1200
GCTTTACTGT ATCTTGCACA TCGGTTTGAC TATAATGATT TATGACTTGT TGATATTTAG 1260

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TGGAAGTAAT ATTGCAAAGT AATATATTTT CTATTATATG TTTATACGAT ATTCGATATT 1320
CCCACCCGTT GTCGCGTTTA CGGAAATACG CCATTGATAT ACTCCACATT AGCTAAAGAA 1380
CAGGGTGTTT AAGGCTACCT TGATGGAAAA GGCTCTCTTA GAGATATTTG TAAATGGTAT 1440
GATATCTCAA GTCGCTCTGT TCTCCAAAAG TGGATAAAAC GGTATACTAG TGGTGAAGAC 1500
TTGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAAG GAAGGCAAGC CACATTTGAA 1560
GAACGTGTAG AGATTGTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 1620
GAGAAGTTTG GTGTTTCCTA CCAACAAATT TATTCTTGGG TCGTAAGCT TGAGAAGAAT 1680
GGCTCACAAG GTTTGGTTGA TAGACGTGTG AAAGGGTTGG AGAGTAGGCC TGATTTAACC 1740
GAGATTGAGC AACTTTAACT CAAGATTAAC CAATTGGAGG AACGTAATCG TCTCTTAGAA 1800
ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 1860
TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 1920
TGCCTATTCA GGCCTTATGC CAACTCTTGA AGGGGTCTCG TTCAGGCTAT TACAAGTGGC 1980
TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 2040
AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100
GTCAACTTGG GACAACTTAA AACAAGAAAC GGATTCGTTG ATTGATGAAC ATTCTGGGGA 2160
TTAGTTCAGT CATTCGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTTACG 2220
AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGACAG 2280
ATGTCACCTA TCTTCAATAC GGTCTGGGAG CTAAAGCTTA TCTCAGTGCG ATTAAAGACC 2340
TGTATAACGG TTCTATTATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTTAT 2400
GAAGACCATT AAAAAGGGGC TAGAGCTCAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460
TTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 2520
CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAAA GTTCTTTGG 2580
GTTTTTCAAG ACTGAGTCTT ACCACCTTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640
TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700
CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760
CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTTCTA AATTGCTAA AATAGCTACA 2820
AGAAAACGAG CCATTTAATG CTTATTCTT ATACTGTCTT GCCTCAGCT CTCCTCGACC 2880
AAAAATTGAG CGTGAGGCTT TTTGTTTCAT TAAACGATGA TATTTCATA TTCATCAGTT 2940
TGTTTTCCGA GAGCCATCAA AGCTTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000
CTAAAAATAA TATATAAAAA AACCTGGCCT ATTTTTCCTG CGTAAAAATT ATGCGCTCCA 3060
ATGCCGCCCA AAAGAACGTT AATAAACAT AAACACTAT GTTAGCATAA GACTTTATTT 3120

TTACAACTGA ATTTTCATATA AATGGATTAG AGTAAGGGAT AAAAGAAATT AGCATAGCTC 3180
TTTTGAAAAT AAAAAAATTA ATATAATATG GAAAAAATTT TATTTTCATAA ACGTTTCATA 3240
AAAGGTATGT AATCTAGTAT TTAGGCAACA CTATTTTGTCT ACTGGTGTCT AGTAACTTAT 3300
AGATTGATAA TTTTACTAGT AAACGTAATT CTTGCTTTA AGAGTTAAAT GTCTATTTAT 3360
TGTAAGCTAA ATTGGGAGGT GAACCTTATGT AAAATTAGAT AGGTACTGTC AAGTACGGGA 3420
TGATTATTGA AACAGCCAGT ATGCATCATA AAATCTGTAT TGCTTAATAA CTATTTCTTT 3480
AACCAGACAT CAGTTCATTG TTTATCATCG CTACCCTAAG TCTAGTTTTT TCAATAGAGC 3540
ATTAGGTAGT TTTTGATAAT AAACTATAT AAACATGAGA ATTAGATTTT GTATTGCATT 3600
CTTCATAATG AGTTATTGA GATTTTCCTT TGAATAAATA GATACGAAAT TCAGTAACTT 3660
CATATATAAA CGGCTCTATC ATTGAGATAG TTTGTCAAAT GAAGAAATTT TTAATGGAAA 3720
TAGTTTTTAAA AACATTAGTT GTAGGCGATG TAAAAATATT AATCCAGTGG ATGCAATAGT 3780
TGCGGAGTAA AAATAGAGAG GAGTAATTAG GAAGTGATAA AAAATGCTAT AGCATATATT 3840
ACCAGAAAAA AAAATAGAAC ACTTATTATA TTTGCTATTT TAACAATTGT TCTTTCTTGC 3900
TTGTATTCAT GTTTAACAAT AATGAAATCA AGTAATGAAA TAGAAAAGGC TTTATATGAA 3960
M K S S N E I E K A L Y E
|---->
AGTTCTAATT CTTCAATATC AATTACAAAA AAAGATGGTA AATATTTTAA TATTAATCAA 4020
S S N S S I S I T K K D G K Y F N I N Q
TTTAAGAATA TTGAAAAAT AAAAGAGGTT GAAGAAAAAA TATTTCAATA TGATGGATTA 4080
F K N I E K I K E V E E K I F Q Y D G L
GCAAAATGA AAGATCTTAA AGTAGTTAGT GGTGAGCAAA GTATAAATAG AGAAGATTTA 4140
A K L K D L K V V S G E Q S I N R E D L
TCTGACGAAT TAAAAATGT TGTTTCACTA GAAGCTACAA GTAATACTAA AAGAAATCTT 4200
S D E F K N V V S L E A T S N T K R N L
TTATTAGTA GTGGAGTATT TAGTTTTTAAA GAAGGAAAAA ATATAGAAGA AAATGATAAG 4260
L F S S G V F S F K E G K N I E E N D K
AATTCAATTC TTGTTTCATGA AGAATTGCT AAACAAAACA AACTAAAATT GGGTGATGAA 4320
N S I L V H E E F A K Q N K L K L G D E
ATTGATCTTG AATTACTAGA TACGGAAAAA AGTGGAAAAA TAAAAAGTCA TAAATTTAAA 4380
I D L E L L D T E K S G K I K S H K F K
ATTATAGGAA TCTTTTCTGG TAAAAACAG GAAACATATA CAGGATTATC ATCTGATTTT 4440
I I G I F S G K K Q E T Y T G L S S D F
AGCGAAAATA TGGTTTTTGT AGATTATTCA ACTAGCCAAG AAATATTAAA TAAATCAGAG 4500
S E N M V F V D Y S T S Q E I L N K S E
AATAATAGAA TTGCAATAA AATTTTAATG TATTCTGGTA GTTTAGAATC TACAGAGCTT 4560
N N R I A N K I L M Y S G S L E S T E L
GCCTTAAACA AATTGAAAGA CTTTAAATTT GATAAGTCAA AGTATTCTAT TAAGAAAGAT 4620

A L N K L K D F K I D K S K Y S I K K D
 AATAAAGCAT TCGAAGAGTC TTTAGAGTCA GTGAGTGGAA TAAAACATAT AATTAAAATA 4680
 N K A F E E S L E S V S G I K H I I K I
 ATGACTTATT CGATTATGTT AGGTGGAATA GTTGTCTTT CATTAACTCTT GATTCTATGG 4740
 M T Y S I M L G G I V V L S L I L I L W
 TTAAGAGAAA GAATTTATGA AATAGGTATA TTTTATCTA TTGGAACAAC TAAGATACAA 4800
 L R E R I Y E I G I F L S I G T T K I Q
 ATTATAAGGC AATTTATATT TGAGTTAATA TTCATATCAA TACCAAGTAT AATATCCTCC 4860
 I I R Q F I F E L I F I S I P S I I S S
 TTATTTTATG GGAATCTACT ATTAAGTA ATTGTAGAAG GATTTATTAA CTCAGAGAAC 4920
 L F L G N L L L K V I V E G F I N S E N
 TCAATGATTT TCGGTGGAAG TTTAATAAAT AAAAGCAGTT TTATGTTAAA CATAACAACA 4980
 S M I F G G S L I N K S S F M L N I T T
 CTTGCAGAAA GTTATTTAAT ATTAATAAGT ATTATTGTTT TATCAGTTGT AATGGCCTCT 5040
 L A E S Y L I L I S I I V L S V V M A S
 TCATTAATAT TATTTAAGAA ACCACAAGAA ATATTATCAA AAATAAGTTA GGAGCAAATA 5100
 S L I L F K K P Q E I L S K I S .
 ATGGATATAT TAGAAATAAA GAATGTAAAT TACAGTTACG CAAATCTAA AGAAAAAGTT 5160
 M D I L E I K N V N Y S Y A N S K E K V
 |---->
 TTGTCAGGAG TAAATCAAAA ATTTGAACCT GGAAAGTTTT ATGCGATAGT AGGGAAGTCA 5220
 L S G V N Q K F E L G K F Y A I V G K S
 GGAACAGGAA AATCCACACT TCTTTCCTTA CTTGCAGGAC TTGATAAAGT TCAAACAGGA 5280
 G T G K S T L L S L L A G L D K V Q T G
 AAAATCTTGT TTAAGAATGA AGATATAGAA AAGAAAGGAT ATAGTAATCA CAGAAAAAAT 5340
 K I L F K N E D I E K K G Y S N H R K N
 AATATATCTT TGGTATTTC AATATATAAT TTAATAGATT ATTTATCGCC GATTGAAAAT 5400
 N I S L V F Q N Y N L I D Y L S P I E N
 ATTAGACTAG TAAATAAATC AGTAGATGAG AGTATCTTGT TCGAATTAGG TTTAGATAAA 5460
 I R L V N K S V D E S I L F E L G L D K
 AAACAAATAA AAAGAAATGT TATGAAATTA TCTGGTGGTC AGCAACAAAG GGTAGCTATT 5520
 K Q I K R N V M K L S G G Q Q Q R V A I
 GCTAGGGCAC TGGTATCAGA TGCCCCAATA ATACTAGCTG ATGAGCCTAC CGGTAACCTA 5580
 A R A L V S D A P I I L A D E P T G N L
 GACAGTGTTA CTGCTGGAGA AATAATT (SEQ ID NO:27) 5607
 D S V T A G E I I .

FIG. 5a

IQSLTEGQLR SDIPEFRAGD TVRVHAKVVE GTRERIQIFE GVVISRKGQG 50
ISEMYTVRKI SGGIGVERTF PIHTPRVDKI EVVRYGKVRK AKLYYLRLAQ 100
GKAARIKEIR R (SEQ ID NO:28) 111

FIG. 5b

MRFAECLGLT VNDIDYTNKY LSINKTWDYH FNQRYLPTKN KSSIRNIPID 50
NDTLFFLHEF TKNKNDRLED KLSNNAVNKT IRKITGREVR VHSLRHTFAS 100
YLISISQVLD HENLNITLEV YAHQLQEOKD RNDKLNQRNL GQNSSKPLFT 150
CNEYVPCRNK TSNYSLGGSC YIH (SEQ ID NO:29) 173

FIG. 5c

MKSSNEIEKA LYESSNSSIS ITKKDGKYFN INQFKNIEKI KEVEEKIFQY 50
DGLAKLKDLD VVSGEQSINR EDLSDEFKNV VSLEATSNTK RNLLFSSGVF 100
SFKEGKNIEE NDKNSILVHE EFAKQNKLLK GDEIDLELLD TEKSGGIKSH 150
KFKIIGIFSG KKQETYTGLS SDFSNNMVV DYSTSQEILN KSENNRIANK 200
ILMYSGSLES TELALNKLKD FKIDKSKYSI KKDKNKAFES LESVSGIKHI 250
IKIMTYSIML GGIVVLSLIL ILWLRERIYE IGIFLSIGTT KIQIIRQFIF 300
ELIFISIPSI ISSLFLGNLL LKVIVEGFIN SENSMIFGGS LINKSSFMLN 350
ITTLAESYLI LISIIVLSV MASSLILEKK PQEILSKIS 389
(SEQ ID NO:30)

FIG. 5d

MDILEIKNVN YSYANSKEKV LSGVNQKFEL GKFYAIVGKS GTGKSTLLSL 50
LAGLDKVQTG KILFKNEDIE KKGYSNHRKN NISLVFQNYN LIDYLSPIEN 100
IRLVNKSVDL SILFELGLDK KQIKRNVMLK SGGQQQRVAI ARALVSDAPI 150
ILADEPTGNL DSVTAGEII (SEQ ID NO:31) 169

FIG. 5e

CATATGACAA TATTTTTC	AA AGTCTACATC ACTTACTCGC	CTGTCTGGA AAATCTGGCA	60
ATACATTAAT CGACCAATTA	GTTGCTGATG GTTTACTTCA	TGCAGATAAT CACTACCATT	120
TTTTC AATGG GAAGTCTCTG	GCCACTTTCA ATACTAACCA	ATTGATTGCG GAAGTTGTCT	180
ATGTTGAAAT ATCCTTAGAT	ACTATGTCTA GTGGTGAACA	TGATTTAGTA AAAGTTAACA	240
TTATCAGACC CACTACCGAG	CATACTATCC CCACGATGAT	GACAGCTAGC CCCTATCATC	300
AAGGTATCAA TGATCCTGCC	GCAGACCAAA AAACATACCA	AATGGAGGGT GCGCTAGCAG	360
TTAAACAGCC TAAACACATA	CAAGTTGACA CAAAACCATT	TAAAGAAGAA GTAAAACATC	420
CTTCAAAATT ACCCATCAGC	CCTGCAACTG AAAGCTTCAC	ACACATTGAC AGTTATAGTC	480
TCAATGACTA TTTTCTTTCT	CGTGGTTTTG CTAATATATA	CGTTTCAGGT GTGGGTACTG	540
CTGGCTCTAC GGGTTTCATG	ACCAGTGGGG ATTACCAACA	AATACAAAGC TTTAAAGCAG	600
TCATTGATTG GTTAAATGGT	AAGGTTACTG CATTCAACAAG	TCATAAACGA GATAAACAAAG	660
TCAAGGCTGA TTGGTCAAAC	GGCCTTGTAG CAACCACAGG	TAAATCTTAT CTCGGTACCA	720
TGTCAACTGG TTTAGCAACA	ACTGGCGTTG AGGGGCTGAA	AGTCATTATC GCTGAAGCCG	780
CAATCTCCAC ATGGTATGAT	TATTATCGAG AAAATGGGCT	TGTGTGTAGT CCAGGCGGCT	840
ACCCCGGTGA AGATTTAGAC	GTTTTAACAG AATTAACATA	CTCACGAAAC CTCTTAGCTG	900
GTGATTACAT CAAAAACAAC	GATTGCTATC AAGCATTGTT	AAATGAACAA TCAAAAGCAA	960
TTGACCGTCA AAGTGGGGAT	TACAACCAAT ACTGGCATGA	CCGTAATTAC CTAATCAGC	1020
TCAATAATGT CAAAAGTCGA	GTAGTTTACA CTCATGGACT	ACAGGATTGG AATGTTAAGC	1080
CAAGACATGT CTACAAAGTT	TTCAATGCAT TGCCTCAAAC	CATCAAAAAA CACCTTTTTT	1140
TACATCAAGG TCAACATGTG	TATATGCATA ATTGGCAGTC	GATTGATTTT CGTGAAAGCA	1200
TGAATGCCTT ACTAAGCCAA	GAAGTACTTG GCATTGACAA	TCATTTCCAA TTAGAAGAGG	1260
TCATTTGGCA AGATAATACT	ACTGAGCAAA CTTGGCAAGT	TTTAGATGCT TTCGGAGGAA	1320
ACCATCAAGA GCAAATTGGT	TTAGGTGATA GTAAAAACT	TATTGATAAC CATTATGACA	1380
AAGAAGCCTT TGATACTTAT	TGTAAAGACT TCAATGTGTT	CAAAAATGAT CTTTTCAAGG	1440
GAAATAATAA AACCAATCAA	ATCACTATTA ATCTTCCTCT	AAAGAAAAAT TATCTCTGTA	1500
ATGGACAGTG CAAACTCCAT	CTACGTGTTA AAAGTACTGA	CAAAAAGGCC ATTTTATCAG	1560
CCCAAATCTT AGACTATGGT	CCTAAAAAAC GATTCAAAGA	TACACCAACC ATCAAATTCT	1620
TAAACAGCCT TGATAATGGT	AAAAATTTTG CCAGAGAAGC	TTTACGTGAA CTCCCGTTTA	1680
CTAAAGATCA TTATCGTGTC	ATCAGTAAAG GTGTCTTGAA	CCTTCAAAAT CGTACAGACT	1740
TACTTACAAT TGAGGCTATC	GAGCCAGAAC AATGGTTTGA	TATCGAGTTT AGCCTCCAAC	1800
CAAGTATATA TCAATTGAGT	AAAGGTGATA ATCTAAGGAT	TATCCTTTAT ACAACTGATT	1860
TTGAACATAC CATTCGAGAT	AATGCTAGTT ACTCTATAAC	AGTAGATTG AGTCAATCTT	1920
ATTTAACTAT CCCAACTAAT	CAAGGAAATT AACTTATGAA	ACTTCTTACT AAAGAACGGT	1980
TTGATGATT CCAACACTTT	TGGTACCAGA TCAATTTATT	ACAAGAGAGT AACTTCGGAG	2040
CAGTTTTTGA CCATGATAAT	AAAAACATTC CACAGGTTGT	TGCAACTATT GTTGATGATT	2100
TACAAGGTT CGGAAGTTTC	AATCATTTCT GGTATTTTGG	CAATACTACT GATACTTCCA	2160
TCCTTATGAT TGCTCATTTA	AATCGAAAAT TCTATATTCA	GGTTAATTTA AAGGACTTTG	2220
ACTTTGCACT CAATTTAATA	GCTATAAATA ATTGGAAGAG	TCTCCTCCAA ACTCAACTTG	2280
AAGCTCTAAA CGATACCCTA	GCAATATTTT AATAAATAAG	GTAGAATGGA GTGACAAAGC	2340
AACGCGAGGG AGACTGATTA	ATGTCATCTT ATTGGAATAA	CTATCCTGAA CTTAAAAAAA	2400

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ATATTGATGA AACCAATCAA CTAATTCAAG AAAGAATACA GGTCAGAAAT AAAGATATTG 2460
AAGCGGCGCT AAGCCAACCTC ACAGCTGCGG GAGGAAAACA GCTCAGACCA GCATTCTTTT 2520
ACCTTTTTTC TCAACTTGGT AATAAGGAGA ATCAAGATAC TCAGCAACTA AAGAAAATCG 2580
CTGCTTCTTT AGAAATCCTT CACGTTGCTA CATTAATCCA TGATGATGTC ATTGATGACT 2640
CACCACCTAAG ACGTGGAAAT ATGACCATTG AAAGCAAGTT TGGCAAAGAC ATCGCAGTTT 2700
ATACTGGGGA TTTACTTTTC ACAGTCTTTT TCGATCTTAT TTTAGAATCT ATGACTGATA 2760
CACCATTTAT GAGGATTAAT GCAAAATCTA TCGGTAAAAT TCTCATGGGA GAATTGGACC 2820
AGATGCACCT TCGTTACAAT CAACAACAAG GTATCCATCA CTATTTACGT GCGATTTTCAG 2880
GTAAGACAGC CGAACTCTTT AAATTAGCTA GCAAAGAAGG AGCTTACTTT GGTGGTGCAG 2940
AGAAGGAGGT TGTTCGTCTA GCAGGCCATA TCGGCTTTAA CATTGGTATG ACATTCCAAA 3000
TTTTGGATGA TATCCTGGAT TATACTGCAG ATAAAAAAC ATTTAATAAG CCTGTCTTAG 3060
AGGATTTAAC ACAAGGCGTT TACAGCCTTC CTCTACTTCT TGCCATTGAA GAAAATCCTG 3120
ATATTTTCAA ACCTATTTTA GATAAAAAAA CAGATATGGC TACTGAAGAC ATGGAAAAAA 3180
TTGCTTATCT CGTCGTTTCC CATAGAGGTG TTGACAAAGC TCGCCATCTA GCTCGTAAAT 3240
TTACTGAGAA AGCTATTAGT GACATAAATA AGCTACCCCA GAACTCTGCA AAAAAACAGT 3300
TGCTACAATT AACTAATTAC CTTTTAAAC GCAAATTTA AATAATAAAA AAACATTCCA 3360
CAATGCTAGA AAAGCAGTTA GGGAAATGTTT TTTTATTATC ATTTATTTAT CGCACCTATC 3420
AATCATCATA GATCACCATC ATCAGCGGCT TTCAGCTGAC GGTAACGTTG ACTACTTTGA 3480
GACAATTCTT GAGGAGAACC TTCCAACCTC AATTGCCCAT TTTCTATAAA TAAGATACGA 3540
TCAGCATGTT CAATACCTTT TAAGTGATGT GTAATCCAAA CTAAGGTCTT ACCTTCCAAT 3600
TCTTTCATAA ATACCCTTAG TAAGGCTTGT TCAGTAATAG GATCAAGTCC AACAGTTGGC 3660
TCATCTAAGA TAACAATTGG GACATCTTTT AGTAAGATTC TAGCCAAAGC AATTCTATGC 3720
CTTTCGCCAC CTGAAAACCT AAGTCCAGCT TCATCAACCA TTGTATAGAG ACCATCTGAT 3780
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CCGCCTAGGG TAATCTTCCC TTGACTTGCT TTCAAGTCGC CACGAAGTAG ACTAGCTAAG 4020
GTACTCTTGC CAGAACCACT CCGCCCTAAA ATAGCAATTT TTTCTCCTTC TTTAATATCC 4080
AAATCTAAAT GATGCAAAAC CCATTTCTCT TGTGGCTTAT ACTGGAAACT TAAATTCCTG 4140
ACGGAAAAAT CATATGGCTT ATTAGGCAAT T (SEQ ID NO:32) 4171

```

FIG. 6a

YDNIFQSLHH LLACRGKSGN TLIDQLVADG LLHADNHYHF FNGKSLATFN 50
 TNQLIREVVY VEISLDTMSS GEHDLVKVNI IRPTTEHTIP TMMTASPYHQ 100
 GINDPAADQK TYQMEGALAV KQPKHIQVDT KPFKEEVKHP SKLPISPATE 150
 SFTHIDSYSL NDYFLSRGFA NIYVSGVGTA GSTGFMTSGD YQQIQSFKAV 200
 IDWLNGKVTA FTSHKRDQV KADWSNGLVA TTGKSYLGTM STGLATTGVE 250
 GLKVIIAEAA ISTWYDYYRE NGLVCSPGGY PGEDLDVLTE LTYSRNLLAG 300
 DYIKNNDCYQ ALLNEQSKAI DRQSGDYNQY WHDRNYLTHV NNVKSRVVYT 350
 HGLQDWNVVK RHVYKVFNAL PQTIKKHLFL HQGQHVYMHN WQSIDFRESM 400
 NALLSQELLG IDNHFQLEEV IWQDNTTEQT WQVLDAFGGN HQEQIGLGDS 450
 KKLIDNHYDK EAFDTYCKDF NVFKNDLEFG NNNKTNQITIN LPLKKNYLLN 500
 GQCKLHLRVK TSDKKAILS A QILDYGPCKR FKDTPTIKFL NSLDNGKNFA 550
 REALRELFFT KDHYRVISKG VLNLNQRTDL LTIEAIEPEQ WFDIEFSLQP 600
 SIYQLSKGDN LRIILYTTDF EHTIRDNASY SITVDLSQSY LTIPTNQGN 649
 (SEQ ID NO:33)

FIG. 6b

MKLLTKERFD DSQHFYQIN LLQESNFGAV FDHDNKNIPQ VVATIVDDLQ 50
 GSGSSNHFWY FGNTTDTISIL MIAHLNRKFY IQVNLKDFDF ALNLIAINNW 100
 KSLLOTQLEA LNDTLAIFQ (SEQ ID NO:34) 119

FIG. 6c

MSSYWNNYPE LKKNIDETNQ LIQERIQVRN KDIEAALSOL TAAGGKQLRP 50
 AFFYLFSQLG NKENQDTQQL KKIAASLEIL HVATLIHDDV IDDSPLRRGN 100
 MTIQSKFGKD IAVYTGDLLE TVFFDLILES MTDTPFMRIN AKSMRKILMG 150
 ELDQMHLRYN QQQGIHHYLR AISGKTAELEF KLASKEGAYF GGAEKEVVRL 200
 AGHIGFNIGM TFQILDDILD YTADKKTFNK PVLEDLTQGV YSLPLLLAIE 250
 ENPDIFKPIL DKKTDMATED MEKIAYLVVS HRGVDKARHL ARKFTEKAIS 300
 DINKLPQNSA KKQLLQLTNY LLKRKI (SEQ ID NO:35) 326

FIG. 6d

LPNKPYDFSV KNLSFQYKPQ EKWVLHHLDL DIKEGEKIAI LGRSGSGKST 50
LASLLRGDLK ASQGKITLGG ADVSIVGDCI SNYIGVIQQA PYLFNTTLLN 100
NIRIGNQDAS EEDVWKVLER VGLKEMVTDL SDGLYTMVDE AGLRFSGGER 150
HRIALARILL KDVPIVILDE PTVGLDPITE QALLRVFMKE LEGKTLVWIT 200
HHLKGIEHAD RILFIENGQL ELEGSPQELS QSSQRYRQLK AADDGDL 247
(SEQ ID NO:36)

FIG. 6e

AATTCTATTT GGAGGTTTTT CTTGAATAAA TGGTTAGTTA AGGCAAGTTC CTTAGTTGTT	60
TTAGGTGGTA TGGTTTTATC TCGGGTTCC CGAGTTTTCG CGGATACTTA TGTCCGTCCA	120
ATTGATAATG GTAGAATTAC AACAGGTTTC AATGGTTATC CTGGACATTG TGGGGTGGAT	180
TATGCTGTTT CGACTGGAAC GATTATTAGG GCAGTGGCAG ATGGTACTGT GAAATTTGCA	240
GGAGCTGGAG CCAACTTTTC TTGGATGACA GACTTAGCAG GAAATTGTGT CATGATTCAA	300
CATGCGGATG GAATGCATAG TGGTTACGCT CATATGTCAC GTGTGGTGGC TAGGACTGGG	360
GAAAAAGTCA AACAAGGAGA TATCATCGGT TACGTAGGAG CAACTGGTAT GGCACGCGGA	420
CCTCACCTTC ATTTTGAATT TTTACCAGCT AACCTAATT TTCAAAATGG TTTCCATGGA	480
CGTATCAATC CAACGTCCT AATTGCTAAC GTTGCACCT TTAGTGGAAA AACGCAAGCA	540
TCAGCTCCAA GCATTAAGCC ATTACAATCA GCTCCTGTAC AGAATCAATC TAGTAAATTA	600
AAAGTGATC GAGTAGATGA ATTACAAAAG GTTAATGGTG TTTGGTTAGT CAAAAATAAC	660
ACCCTAACGC CGACTGGGT TGAATGGAAC GATAATGGTA TACCAGCATC AGAAATTGAT	720
GAGGTTGATG CTAATGGTAA TTTGACAGCT GACCAGGTTT TTCAAAAAGG TGGTTACTTT	780
ATCTTTAATC CTAAACTCT TAAGACTGTA GAAAAACCCA TCCAAGGAAC AGCTGGTTTA	840
ACTTGGGCTA AGACACGCTT TGCTAATGGT AGTTCAGTTT GGCTTCGCGT TGACAACAGT	900
CAAGAACTGC TTTACAAATA GTTTGAGGTA TTGATTCATT GTTTTAAATG ACAGTTTTGT	960
TACTAACTAA GTACAATTC TTTAAACCGT CTGAAAATAA TTTTATAGTC CAGTAAAGTG	1020
TGATATTATA GTCTCGGACT AATAAAAAGG AAATAGGAAT TGAAGCAATG AAAATGAATA	1080
AAAAGGTACT ATTGACATCG ACAATGGCAG CTTGCTATT ATCAGTCGCA AGTGTTCAG	1140
CACAAGAAAC AGATACGACG TGGACAGCAC GTACTGTTTC AGAGGTAAAG GCTGATTTGG	1200
TAAAGCAAGA CAATAAATCA TCATATACTG TGAAATATGG TGATACACTA AGCGTTATTT	1260
CAGAAGCAAT GTCAATTGAT ATGAATGTCT TAGCAAAAAT TAATAACATT GCAGATATCA	1320
ATCTTATTTA TCCTGAGACA AACTGACAG TAACTTACGA TCAGAAGAGT CATACTGCCA	1380
CTTCAATGAA AATAGAAACA CCAGCAACAA ATGCTGCTGG TCAAACAACA GCTACTGTGG	1440
ATTTGAAAAC CAATCAAGTT TCTGTTGCAG ACCAAAAAGT TTCTCTCAAT ACAATTTCCG	1500
AAGGTATGAC ACCAGAAGCA GCAACAACGA TTGTTTCGCC AATGAAGACA TATTCCTCTG	1560
CGCCAGCTTT GAAATCAAAA GAAGTATTAG CACAAGAGCA AGCTGTTAGT CAAGCAGCAG	1620
CTAATGAACA GGTATCAACA GCTCCTGTGA AGTCGATTAC TTCAGAAGTT CCAGCAGCTA	1680
AAGAGGAAGT TAAACCAACT CAGACGTCAG TCAGTCAGTC AACACAGTA TCACCAGCTT	1740
CTGTTGCCGC TGAAACACCA GCTCCAGTAG CTAAAGTAGC ACCGGTAAGA ACTGTAGCAG	1800
CCCCTAGAGT GGCAAGTGTT AAAGTAGTCA CTCCTAAAGT AGAACTGGT GCATCACCAG	1860
AGCATGTATC AGCTCCAGCA GTTCTGTGA CTACGACTTC AACAGCTACA GACAGTAAAGT	1920
TACAAGCGAC TGAAGTTAAG AGCGTTCCGG TAGCACAAAAGCTCCAACA GCAACACCGG	1980
TAGCACAAAC AGCTTCAACA ACAAATGCAG TAGCTGCACA TCCTGAAAAT GCAGGGCTCC	2040
AACCTCATGT TGCAGCTTAT AAAGAAAAAG TAGCGTCAAC TTATGGAGTT AATGAATTC	2100
GTACATACCG TGCAGGTGAT CCAGGTGATC ATGGTAAAGG TTTAGCAGTC GACTTTATTG	2160
TAGGTAAAAA CCAAGCACTT GGTAATGAAG TTGCACAGTA CTCTACACAA AATATGGCAG	2220
CAAATAACAT TTCATATGTT ATCTGGCAAC AAAAGTTTTA CTCAAATACA AATAGTATTT	2280
ATGGACCTGC TAATACTTGG AATGCAATGC CAGATCGTGG TGGCGTTACT GCCAACCATT	2340
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TAAATCTAA GTCTGTAAAG ATTATTGAAA ACGCTTTAAA AACAGATATA ATAAGGTTTG 2580
TAGATATCTA AAATTAATAA AGATAAGGAA GTGAGAATAT GCCACATCTA AGTAAAGAAG 2640
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GTACACTTAG AGAGCGTCAT GATGTTTGA GTGTAGCTGA GTTTATTCAA AAGATAAAAG 3000
GGAAATATCC TGAACAGTTG CTAATGGCTG ATATAAGTAC TTTTGAAGAA GGTAAAAATG 3060
CTTTTGAAGC AGGAGTTGAT TTTGTGGGA CAACTCTATC TGGATACACA GATTACAGCC 3120
GCCAAGAAGA AGGACCGGAT ATAGAACTCC TTAATAAGCT TTGTCAAGCC GGTATAGATG 3180
TGATTGCGGA AGGTAAAT CATACTCCTA AGCAAGCTAA TGAAATTAAT CATATAGGTG 3240
TTGCAGGAAT TGTAGTTGGT GGTGCTATCA CTAGACCAA AGAAATAGCG GAGCGTTTCA 3300
TCTCAGGACT TAGTTAAAAG TGTTACTCAA AAATCAAAT CAAAATAAAA AAGGGGAATA 3360
GTTATGAGTA TCAAAAAAAG TGTGATTGGT TTTTGCCCTG GAGCTGCAGC ATTATCAATG 3420
TTTGCTTGTG TAGACAGTAG TCAATCTGTT ATGGCTGCCG AGAAGGATAA AGTCGAAATT 3480

```

(SEQ ID NO:37)

FIG. 7a

```

NSIWRFFLNK WLVKASSLV LGGMVL SAGS RVLADTYVRP IDNGRITTGF 50
NGYPGHCGVD YAVPTGTIIR AVADGTVKFA GAGANFSWMT DLAGNCVMIQ 100
HADGMHSGYA HMSRVVARTG EKVKQGDII G YVGATGMATG PHLHFEFLPA 150
NPNFQNGFHG RINPTSLIAN VATFSGKTQA SAPSIKPLQS APVQNQSSKL 200
KVYRVDELQK VNGVWLKNN TLTPTGFDWN DNGIPASEID EVDANGNLTA 250
DQVLQKGGYF IFNPKTLKTV EKPIQGTAGL TWAKTRFANG SSVWLRVDNS 300
QELLYK (SEQ ID NO:38) 306

```

FIG. 7b

MKMNKKVLLT	STMAASLLSV	ASVQAQETDT	TWTARTVSEV	KADLVKQDNK	50
SSYTVKYGDT	LSVISEAMSI	DMNVLAKINN	IADINLIYPE	TTLTVTYDQK	100
SHTATSMKIE	TPATNAAGQT	TATVDLKTNO	VSVADQKVSL	NTISEGMTPE	150
AATTIVSPMK	TYSSAPALKS	KEVLAQEQAV	SQAAANEQVS	TAPVKSITSE	200
VPAAKEEVKP	TQTSVSQSTT	VSPASVAAET	PAPVAKVAPV	RTVAAPRVAS	250
VKVVT PKVET	GASPEHVSAP	AVPVTTTSTA	TDSKLQATEV	KSVPVAQKAP	300
TATPVAQPAS	TTNAVAAHPE	NAGLQPHVAA	YKEKVASTYG	VNEFSTYRAG	350
DPGDHGKGLA	VDFIVGKNQA	LGNEVAQYST	QNMAANNISY	VIWQQKFYSN	400
TNSIYGPANT	WNAMPDRGGV	TANHVDHVHV	SFNK	(SEQ ID NO:39)	434

FIG. 7c

MPHLSKEAFK	KQIKNGIIVS	CQALPGEPLY	TESGGVMPLL	ALAAQEAGAV	50
GIRANSVRDI	KEIQEVTNLP	IIGIIKREYP	PQEPFITATM	TEVDQLASLD	100
IAVIALDCTL	RERHDGLSVA	EFIQKIKGKY	PEQLLMADIS	TFEEGKNAFE	150
AGVDFVGTTL	SGYTDYXRQE	EGPDIELLNK	LCQAGIDVIA	EGKIHTPKQA	200
NEINHIGVAG	IVVGGAITRP	KEIAERFISG	LS	(SEQ ID NO:40)	232

FIG. 7d

MSIKKSVIGF	CLGAAALSMF	ACVDSSQSVM	AAEKDKVEI	39
(SEQ ID NO:41)				

FIG. 7e

ATGAAAATGA	ATAAAAAGGT	ACTATTGACA	TCGACAATGG	CAGCTTCGCT	50
ATTATCAGTC	GCAAGTGTTT	AAGCACAAGA	AACAGATACG	ACGTGGACAG	100
CACGTACTGT	TTCAGAGGTA	AAGGCTGATT	TGGTAAAGCA	AGACAATAAA	150
TCATCATATA	CTGTGAAATA	TGGTGATACA	CTAAGCGTTA	TTTCAGAAGC	200
AATGTCAATT	GATATGAATG	TCTTAGCAAA	AATTAATAAC	ATTGCAGATA	250
TCAATCTTAT	TTATCCTGAG	ACAACACTGA	CAGTAACTTA	CGATCAGAAG	300
AGTCATACTG	CCACTTCAAT	GAAAATAGAA	ACACCAGCAA	CAAATGCTGC	350
TGGTCAAACA	ACAGCTACTG	TGGATTTGAA	AACCAATCAA	GTTTCTGTTG	400
CAGACCAAAA	AGTTTCTCTC	AATACAATTT	CGGAAGGTAT	GACACCAGAA	450
GCAGCAACAA	CGATTGTTTC	GCCAATGAAG	ACATATTCTT	CTGCGCCAGC	500
TTTGAAATCA	AAAGAAGTAT	TAGCACAAGA	GCAAGCTGTT	AGTCAAGCAG	550
CAGCTAATGA	ACAGGTATCA	ACAGCTCCTG	TGAAGTCGAT	TACTTCAGAA	600
GTTCCAGCAG	CTAAAGAGGA	AGTTAAACCA	ACTCAGACGT	CAGTCAGTCA	650
GTCAACAACA	GTATCACCAG	CTTCTGTTGC	CGCTGAAACA	CCAGCTCCAG	700
TAGCTAAAGT	AGCACCGGTA	AGAACTGTAG	CAGCCCCTAG	AGTGGCAAGT	750
GTTAAAGTAG	TCACTCCTAA	AGTAGAAACT	GGTGCATCAC	CAGAGCATGT	800
ATCAGCTCCA	GCAGTTCCTG	TGACTACGAC	TTCAACAGCT	ACAGACAGTA	850
AGTTACAAGC	GACTGAAGTT	AAGAGCGTTC	CGGTAGCACA	AAAAGCTCCA	900
ACAGCAACAC	CGGTAGCACA	ACCAGCTTCA	ACAACAAATG	CAGTAGCTGC	950
ACATCCTGAA	AATGCAGGGC	TCCAACCTCA	TGTTGCAGCT	TATAAAGAAA	1000
AAGTAGCGTC	AACTTATGGA	GTTAATGAAT	TCAGTACATA	CCGTGCAGGT	1050
GATCCAGGTG	ATCATGGTAA	AGGTTTAGCA	GTCGACTTTA	TTGTAGGTAA	1100
AAACCAAGCA	CTTGGTAATG	AAGTTGCACA	GTA CTCTACA	CAAAATATGG	1150
CAGCAAATAA	CATTTTCATAT	GTTATCTGGC	AACAAAAGTT	TTACTCAAAT	1200
ACAAATAGTA	TTTATGGACC	TGCTAATACT	TGGAATGCAA	TGCCAGATCG	1250
TGGTGGCGTT	ACTGCCAACC	ATTATGACCA	TGTTACGTA	TCATTTAACA	1300
AATAA					1305

(SEQ ID NO:42)

FIG. 8

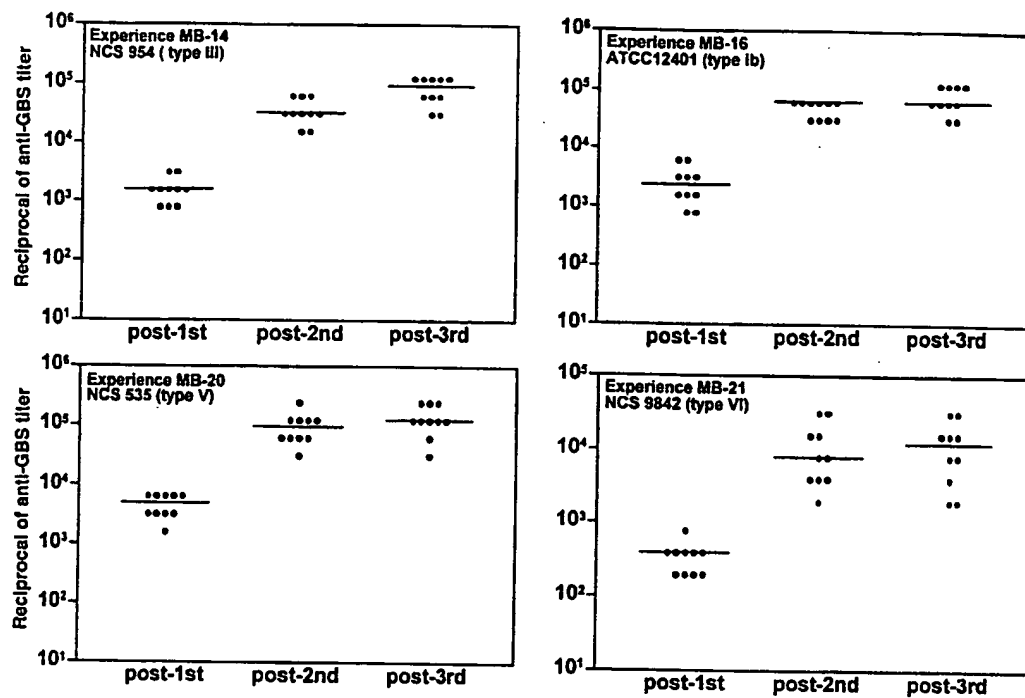
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ATACACTAAG	CGTTATTTCA	GAAGCAATGT	CAATTGATAT	GAATGTCTTA	150
GCAAAAATTA	ATAACATTGC	AGATATCAAT	CTTATTTATC	CTGAGACAAC	200
ACTGACAGTA	ACTTACGATC	AGAAGAGTCA	TACTGCCACT	TCAATGAAAA	250
TAGAAACACC	AGCAACAAAT	GCTGCTGGTC	AAACAACAGC	TACTGTGGAT	300
TTGAAAACCA	ATCAAGTTTC	TGTTGCAGAC	CAAAAAGTTT	CTCTCAATAC	350
AATTTTCGGAA	GGTATGACAC	CAGAAGCAGC	AACAACGATT	GTTTCGCCAA	400
TGAAGACATA	TTCTTCTGCG	CCAGCTTTGA	AATCAAAAGA	AGTATTAGCA	450
CAAGAGCAAG	CTGTTAGTCA	AGCAGCAGCT	AATGAACAGG	TATCAACAGC	500
TCCTGTGAAG	TCGATTACTT	CAGAAGTTCC	AGCAGCTAAA	GAGGAAGTTA	550
AACCAACTCA	GACGTCAGTC	AGTCAGTCAA	CAACAGTATC	ACCAGCTTCT	600
GTTGCCGCTG	AAACACCAGC	TCCAGTAGCT	AAAGTAGCAC	CGGTAAGAAC	650
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ACGACTTCAA	CAGCTACAGA	CAGTAAGTTA	CAAGCGACTG	AAGTTAAGAG	800
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CCTCATGTTG	CAGCTTATAA	AGAAAAAGTA	GCGTCAACTT	ATGGAGTTAA	950
TGAATTCAGT	ACATACCGTG	CAGGTGATCC	AGGTGATCAT	GGTAAAGGTT	1000
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GCACAGTACT	CTACACAAAA	TATGGCAGCA	AATAACATTT	CATATGTTAT	1100
CTGGCAACAA	AAGTTTTACT	CAAATACAAA	TAGTATTTAT	GGACCTGCTA	1150
ATACTTGGA	TGCAATGCCA	GATCGTGGTG	GCGTTACTGC	CAACCATTAT	1200
GACCATGTTC	ACGTATCATT	TAACAAATAA	(SEQ ID NO:43)		1230

FIG. 9

QETDTTWTAR	TVSEVKADLV	KQDNKSSYTV	KYGDTLISVIS	EAMSIDMNVL	50
AKINNIADIN	LIYPETTLTV	TYDQKSHTAT	SMKIETPATN	AAGQTTATVD	100
LKTNQVSVAD	QKVSNTISE	GMTPEAATTI	VSPMKTYSSA	PALKSKEVLA	150
QEQAVSQAAA	NEQVSTAPVK	SITSEVPAAK	EEVKPTQTSV	SQSTTVSPAS	200
VAAETPAPVA	KVAPVRTVAA	PRVASVKVVT	PKVETGASPE	HVSAPAVPVT	250
TTSTATDSKL	QATEVKSVPV	AQKAPTATPV	AQPASTTNAV	AAHPENAGLQ	300
PHVAAYKEKV	ASTYGVNEFS	TYRAGDPGDH	GKGLAVDFIV	GKNQALGNEV	350
AQYSTQNMAA	NNISYVIWQQ	KFYSNTNSIY	GPANTWNAMP	DRGGVTANHY	400
DHVHVSFNK	(SEQ ID NO:44)				409

FIG. 9a

Fig. 10



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SEQUENCE LISTING

<110> BioChem Vaccins
 RIOUX, Clément
 DENIS, Martin
 BRODEUR, Bernard R.
 HAMEL, Josée
 CHARLEBOIS, Isabelle
 BOYER, Martine

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47

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ttt agt atg tct aaa gaa gag ttg tca tat tta ccc gtt att aaa ctt Phe Ser Met Ser Lys Glu Glu Leu Ser Tyr Leu Pro Val Ile Lys Leu 190 195 200 205	686
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atc aat gta ttg cta gtt gct att tat ggt gct ttg aca gtt gat aaa Ile Asn Val Leu Leu Val Ala Ile Tyr Gly Ala Leu Thr Val Asp Lys 240 245 250	830
aaa atc tta tta aaa cag ggt ggt tta cct ata tta gct ctt tta aca Lys Ile Leu Leu Lys Gln Gly Gly Leu Pro Ile Leu Ala Leu Leu Thr 255 260 265	878
ttc tta ttt taatactact tagccgttcg atttagttga acggcctttta Phe Leu Phe 270	927
gtaatcattt ttttctcata atacaggtag ttttaagtaat ttgtcttttaa aaatagtata atataactac gaattcaaag agaggtgact ttgatt atg act gag aac tgg tta Met Thr Glu Asn Trp Leu 275	987 1041
cat act aaa gat ggt tca gat att tat tat cgt gtc gtt ggt caa ggt His Thr Lys Asp Gly Ser Asp Ile Tyr Tyr Arg Val Val Gly Gln Gly 280 285 290	1089
caa ccg att gtt ttt tta cat ggc aat agc tta agt agt cgc tat ttt Gln Pro Ile Val Phe Leu His Gly Asn Ser Leu Ser Ser Arg Tyr Phe 295 300 305 310	1137
gat aag caa ata gca tat ttt tct aag tat tac caa gtt att gtt atg Asp Lys Gln Ile Ala Tyr Phe Ser Lys Tyr Tyr Gln Val Ile Val Met 315 320 325	1185
gat agt aga ggg cat ggc aaa agt cat gca aag cta aat acc att agt Asp Ser Arg Gly His Gly Lys Ser His Ala Lys Leu Asn Thr Ile Ser 330 335 340	1233
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Leu Lys Ile Ser Pro Ala Asp Leu Gln His Val Ser Thr Pro Val Met	
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Val Leu Val Gly Asn Lys Asp Ile Ile Lys Leu Asn His Ser Lys Lys	
455 460 465 470	
ctt gct tct tat ttt cca agg ggg gag ttt tat tct tta gtt ggc ttt	1665
Leu Ala Ser Tyr Phe Pro Arg Gly Glu Phe Tyr Ser Leu Val Gly Phe	
475 480 485	
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Gly His His Ile Ile Lys Gln Asp Ser His Val Phe Asn Ile Ile Ala	
490 495 500	
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Lys Lys Phe Ile Asn Asp Thr Leu Lys Gly Glu Ile Val Glu Lys Ala	
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Asn *	
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Glu Gln Leu Lys Ser Val Phe Gly Gln Leu Ser Pro Met Asn Leu Phe	
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Leu Ile Ile Leu Val Gly Val Ile Ala Val Leu Pro Thr Thr Gly Tyr	
550 555 560	
gac ttt gta ctg aat gga ctt tta cgt aca gat aaa agc aaa agg tat	2014
Asp Phe Val Leu Asn Gly Leu Leu Arg Thr Asp Lys Ser Lys Arg Tyr	
565 570 575	
att tta cag act agt tgg tgt atc aac act ttt aat aac ttg tca gga	2062
Ile Leu Gln Thr Ser Trp Cys Ile Asn Thr Phe Asn Asn Leu Ser Gly	
580 585 590	
ttc ggt ggc tta atc gat att ggg ttg cgc atg gct ttt tat ggt aaa	2110
Phe Gly Gly Leu Ile Asp Ile Gly Leu Arg Met Ala Phe Tyr Gly Lys	
595 600 605	
aaa ggt caa gag aag agt gac cta aga gaa gtg act cgt ttt tta ccc	2158
Lys Gly Gln Glu Lys Ser Asp Leu Arg Glu Val Thr Arg Phe Leu Pro	
610 615 620 625	

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tta att ggt gct agt atg tat ttt cct gtt att tat tgg att tct ggt Leu Ile Gly Ala Ser Met Tyr Phe Pro Val Ile Tyr Trp Ile Ser Gly 660 665 670	2302
cat aaa gga agc cat tat ttc gga gat atg cca tct agt act cgt ata His Lys Gly Ser His Tyr Phe Gly Asp Met Pro Ser Ser Thr Arg Ile 675 680 685	2350
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att ccc ggt gga tta gga agt ttt gaa tta gtt cta ttt aca ggg ttt Ile Pro Gly Gly Leu Gly Ser Phe Glu Leu Val Leu Phe Thr Gly Phe 740 745 750	2542
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aat att act tat att atg tgg ttg cag aag cta ggc ttg gac cca tta Asn Ile Thr Tyr Ile Met Trp Leu Gln Lys Leu Gly Leu Asp Pro Leu 835 840 845	2830

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ttt att ctc tta gct aga act att gat caa aaa gtg aaa aat gct ttt Phe Ile Leu Leu Ala Arg Thr Ile Asp Gln Lys Val Lys Asn Ala Phe 870 875 880	2926
cca att gct att atc tgg att act ttg aca ttg ttt tat ctt aat tta Pro Ile Ala Ile Ile Trp Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu 885 890 895	2974
ggg cat att agt tgg cga cta tct ttc tgg ttt att tta cta ttg tta Gly His Ile Ser Trp Arg Leu Ser Phe Trp Phe Ile Leu Leu Leu 900 905 910	3022
ggc tta tta gtc att aag cca act ctc tat aaa aaa caa ttt att tat Gly Leu Leu Val Ile Lys Pro Thr Leu Tyr Lys Lys Gln Phe Ile Tyr 915 920 925	3070
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atc tct att gat tta atg cgt cac gat aaa cag aaa att ccg aat ggc Ile Ser Ile Asp Leu Met Arg His Asp Lys Gln Lys Ile Pro Asn Gly 1220 1225 1230	3982
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gga tac cac tat ttt gat ttg ggg atg gca cct tta tca gga gtt ggt Gly Tyr His Tyr Phe Asp Leu Gly Met Ala Pro Leu Ser Gly Val Gly 1250 1255 1260 1265	4078
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Ala	Val	Gln 35	Phe	Ile	Gly	Leu	Lys 40	Pro	Asp	Tyr	Pro	Gly 45	Lys	Thr	Tyr
Phe 50	Ile	Ile	Leu	Leu	Thr 55	Ala	Trp	Thr	Leu	Met	Ala 60	Leu	Val	Thr	Ala
Leu 65	Val	Gly	Trp	Asp 70	Asn	Arg	Tyr	Gly	Ser	Phe 75	Leu	Ser	Leu	Leu	Ile 80
Leu	Leu	Phe	Gln 85	Leu	Gly	Ser	Ser	Ala 90	Gly	Thr	Tyr	Pro	Ile 95	Glu	Leu
Ser	Pro	Lys 100	Phe	Gln	Thr	Ile	Gln 105	Pro	Phe	Leu	Pro	Met 110	Thr	Tyr	
Ser	Val 115	Ser	Gly	Leu	Arg	Glu	Thr 120	Ile	Ser	Leu	Thr	Gly 125	Asp	Val	Asn
His 130	Gln	Trp	Arg	Met	Leu	Val 135	Ile	Phe	Leu	Val	Ser 140	Ser	Met	Ile	Leu
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20 25 30

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Gly Lys Ile Phe Ser Met Ser Lys Glu Glu Leu Ser Tyr Leu Pro Val
 35 40 45
 Ile Lys Leu Phe Lys Asn Gln Gly Val Tyr Asn Gly Leu Ile Gly Leu
 50 55 60
 Phe Leu Leu Tyr Gly Leu Tyr Ile Ser Gln Asn Gln Glu Ile Val Ala
 65 70 75 80
 Val Phe Leu Ile Asn Val Leu Leu Val Ala Ile Tyr Gly Ala Leu Thr
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 Val Asp Lys Lys Ile Leu Leu Lys Gln Gly Gly Leu Pro Ile Leu Ala
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 35 40 45
 Tyr Gln Val Ile Val Met Asp Ser Arg Gly His Gly Lys Ser His Ala
 50 55 60
 Lys Leu Asn Thr Ile Ser Phe Arg Gln Ile Ala Val Asp Leu Lys Asp
 65 70 75 80
 Ile Leu Val His Leu Glu Ile Asp Lys Val Ile Leu Val Gly His Ser
 85 90 95
 Asp Gly Ala Asn Leu Ala Leu Val Phe Gln Thr Met Phe Pro Gly Met
 100 105 110
 Val Arg Gly Leu Leu Leu Asn Ser Gly Asn Leu Thr Ile His Gly Gln
 115 120 125
 Arg Trp Trp Asp Ile Leu Leu Val Arg Ile Ala Tyr Lys Phe Leu His
 130 135 140
 Tyr Leu Gly Lys Leu Phe Pro Tyr Met Arg Gln Lys Ala Gln Val Ile
 145 150 155 160
 Ser Leu Met Leu Glu Asp Leu Lys Ile Ser Pro Ala Asp Leu Gln His
 165 170 175
 Val Ser Thr Pro Val Met Val Leu Val Gly Asn Lys Asp Ile Ile Lys
 180 185 190
 Leu Asn His Ser Lys Lys Leu Ala Ser Tyr Phe Pro Arg Gly Glu Phe
 195 200 205
 Tyr Ser Leu Val Gly Phe Gly His His Ile Ile Lys Gln Asp Ser His
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 Glu Ile Val Glu Lys Ala Asn
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Val Ile Ala Val Leu Pro Thr Thr Gly Tyr Asp Phe Val Leu Asn Gly
35 40 45
Leu Leu Arg Thr Asp Lys Ser Lys Arg Tyr Ile Leu Gln Thr Ser Trp
50 55 60
Cys Ile Asn Thr Phe Asn Asn Leu Ser Gly Phe Gly Gly Leu Ile Asp
65 70 75 80
Ile Gly Leu Arg Met Ala Phe Tyr Gly Lys Lys Gly Gln Glu Lys Ser
85 90 95
Asp Leu Arg Glu Val Thr Arg Phe Leu Pro Tyr Leu Ile Ser Gly Leu
100 105 110
Ser Phe Ile Ser Val Ile Ala Leu Ile Met Ser His Ile Phe His Ala
115 120 125
Lys Ala Ser Val Asp Tyr Tyr Tyr Leu Val Leu Ile Gly Ala Ser Met
130 135 140
Tyr Phe Pro Val Ile Tyr Trp Ile Ser Gly His Lys Gly Ser His Tyr
145 150 155 160
Phe Gly Asp Met Pro Ser Ser Thr Arg Ile Lys Leu Gly Val Val Ser
165 170 175
Phe Phe Glu Trp Gly Cys Ala Ala Ala Phe Ile Ile Ile Gly Tyr
180 185 190
Leu Met Gly Ile His Leu Pro Val Tyr Lys Ile Leu Pro Leu Phe Cys
195 200 205
Ile Gly Cys Ala Val Gly Ile Val Ser Leu Ile Pro Gly Gly Leu Gly
210 215 220
Ser Phe Glu Leu Val Leu Phe Thr Gly Phe Ala Ala Glu Gly Leu Pro
225 230 235 240
Lys Glu Thr Val Val Ala Trp Leu Leu Leu Tyr Arg Leu Ala Tyr Tyr
245 250 255
Ile Ile Pro Phe Phe Ala Gly Ile Tyr Phe Phe Ile His Tyr Leu Gly
260 265 270
Ser Gln Ile Asn Gln Arg Tyr Glu Asn Val Pro Lys Glu Leu Val Ser
275 280 285
Thr Val Leu Gln Thr Met Val Ser His Leu Met Arg Ile Leu Gly Ala
290 295 300
Phe Leu Ile Phe Ser Thr Ala Phe Phe Glu Asn Ile Thr Tyr Ile Met
305 310 315 320
Trp Leu Gln Lys Leu Gly Leu Asp Pro Leu Gln Glu Gln Met Leu Trp
325 330 335
Gln Phe Pro Gly Leu Leu Leu Gly Val Cys Phe Ile Leu Leu Ala Arg
340 345 350
Thr Ile Asp Gln Lys Val Lys Asn Ala Phe Pro Ile Ala Ile Ile Trp
355 360 365
Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu Gly His Ile Ser Trp Arg
370 375 380
Leu Ser Phe Trp Phe Ile Leu Leu Leu Leu Gly Leu Leu Val Ile Lys
385 390 395 400
Pro Thr Leu Tyr Lys Lys Gln Phe Ile Tyr Ser Trp Glu Glu Arg Ile
405 410 415
Lys Asp Gly Ile Ile Ile Val Ser Leu Met Gly Val Leu Phe Tyr Ile
420 425 430

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Ala Gly Leu Leu Phe Pro Ile Arg Ala His Ile Thr Gly Gly Ser Ile
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 Glu Arg Leu His Tyr Ile Ile Ala Trp Glu Pro Ile Ala Leu Ala Thr
 450 455 460
 Leu Ile Leu Thr Leu Val Tyr Leu Cys Leu Val Lys Ile Leu Gln Gly
 465 470 475 480
 Lys Ser Cys Gln Ile Gly Asp Val Phe Asn Val Asp Arg Tyr Lys Lys
 485 490 495
 Leu Leu Gln Ala Tyr Gly Gly Ser Ser Asp Ser Gly Leu Ala Phe Leu
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 Asn Asp Lys Arg Leu Tyr Trp Tyr Gln Lys Asn Gly Glu Asp Cys Val
 515 520 525
 Ala Phe Gln Phe Val Ile Val Asn Asn Lys Cys Leu Ile Met Gly Glu
 530 535 540
 Pro Ala Gly Asp Asp Thr Tyr Ile Arg Glu Ala Ile Glu Ser Phe Ile
 545 550 555 560
 Asp Asp Ala Asp Lys Leu Asp Tyr Asp Leu Val Phe Tyr Ser Ile Gly
 565 570 575
 Gln Lys Leu Thr Leu Leu Leu His Glu Tyr Gly Phe Asp Phe Met Lys
 580 585 590
 Val Gly Glu Asp Ala Leu Val Asn Leu Glu Thr Phe Thr Leu Lys Gly
 595 600 605
 Asn Lys Tyr Lys Pro Phe Arg Asn Ala Leu Asn Arg Val Glu Lys Asp
 610 615 620
 Gly Phe Tyr Phe Glu Val Val Gln Ser Pro His Ser Gln Glu Leu Leu
 625 630 635 640
 Asn Ser Leu Glu Glu Ile Ser Asn Thr Trp Leu Glu Gly Arg Pro Glu
 645 650 655
 Lys Gly Phe Ser Leu Gly Tyr Phe Asn Lys Asp Tyr Phe Gln Gln Ala
 660 665 670
 Pro Ile Ala Leu Val Lys Asn Ala Glu His Glu Val Val Ala Phe Ala
 675 680 685
 Asn Ile Met Pro Asn Tyr Glu Lys Ser Ile Ile Ser Ile Asp Leu Met
 690 695 700
 Arg His Asp Lys Gln Lys Ile Pro Asn Gly Val Met Asp Phe Leu Phe
 705 710 715 720
 Leu Ser Leu Phe Ser Tyr Tyr Gln Glu Lys Gly Tyr His Tyr Phe Asp
 725 730 735
 Leu Gly Met Ala Pro Leu Ser Gly Val Gly Arg Val Glu Thr Ser Phe
 740 745 750
 Ala Lys Glu Arg Met Ala Tyr Leu Val Tyr His Phe Gly Ser His Phe
 755 760 765
 Tyr Ser Phe Asn Gly Leu His Lys Tyr Lys Lys Lys Phe Thr Pro Leu
 770 775 780
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 Gln Glu Gln Met Leu Trp Gln Phe Pro Gly Leu Leu Leu Gly Val Cys
 35 40 45
 Phe Ile Leu Leu Ala Arg Thr Ile Asp Gln Lys Val Lys Asn Ala Phe
 50 55 60
 Pro Ile Ala Ile Ile Trp Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu
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 Gly His Ile Ser Trp Arg Leu Ser Phe Trp Phe Ile Leu Leu Leu Leu
 85 90 95
 Gly Leu Leu Val Ile Lys Pro Thr Leu Tyr Lys Lys Gln Phe Ile Tyr
 100 105 110
 Ser Trp Glu Arg Ile Lys Asp Gly Ile Ile Ile Val Ser Leu Met
 115 120 125
 Gly Val Leu Phe Tyr Ile Ala Gly Leu Leu Phe Pro Ile Arg Ala His
 130 135 140
 Ile Thr Gly Gly Ser Ile Glu Arg Leu His Tyr Ile Ile Ala Trp Glu
 145 150 155 160
 Pro Ile Ala Leu Ala Thr Leu Ile Leu Thr Leu Val Tyr Leu Cys Leu
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 Val Lys Ile Leu Gln Gly Lys Ser Cys Gln Ile Gly Asp Val Phe Asn
 180 185 190
 Val Asp Arg Tyr Lys Lys Leu Leu Gln Ala Tyr Gly Gly Ser Ser Asp
 195 200 205
 Ser Gly Leu Ala Phe Leu Asn Asp Lys Arg Leu Tyr Trp Tyr Gln Lys
 210 215 220
 Asn Gly Glu Asp Cys Val Ala Phe Gln Phe Val Ile Val Asn Asn Lys
 225 230 235 240
 Cys Leu Ile Met Gly Glu Pro Ala Gly Asp Asp Thr Tyr Ile Arg Glu
 245 250 255
 Ala Ile Glu Ser Phe Ile Asp Asp Ala Asp Lys Leu Asp Tyr Asp Leu
 260 265 270
 Val Phe Tyr Ser Ile Gly Gln Lys Leu Thr Leu Leu Leu His Glu Tyr
 275 280 285
 Gly Phe Asp Phe Met Lys Val Gly Glu Asp Ala Leu Val Asn Leu Glu
 290 295 300
 Thr Phe Thr Leu Lys Gly Asn Lys Tyr Lys Pro Phe Arg Asn Ala Leu
 305 310 315 320
 Asn Arg Val Glu Lys Asp Gly Phe Tyr Phe Glu Val Val Gln Ser Pro
 325 330 335
 His Ser Gln Glu Leu Leu Asn Ser Leu Glu Glu Ile Ser Asn Thr Trp
 340 345 350
 Leu Glu Gly Arg Pro Glu Lys Gly Phe Ser Leu Gly Tyr Phe Asn Lys
 355 360 365
 Asp Tyr Phe Gln Gln Ala Pro Ile Ala Leu Val Lys Asn Ala Glu His
 370 375 380
 Glu Val Val Ala Phe Ala Asn Ile Met Pro Asn Tyr Glu Lys Ser Ile
 385 390 395 400
 Ile Ser Ile Asp Leu Met Arg His Asp Lys Gln Lys Ile Pro Asn Gly
 405 410 415
 Val Met Asp Phe Leu Phe Leu Ser Leu Phe Ser Tyr Tyr Gln Glu Lys
 420 425 430
 Gly Tyr His Tyr Phe Asp Leu Gly Met Ala Pro Leu Ser Gly Val Gly
 435 440 445

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Arg Val Glu Thr Ser Phe Ala Lys Glu Arg Met Ala Tyr Leu Val Tyr
 450 455 460
 His Phe Gly Ser His Phe Tyr Ser Phe Asn Gly Leu His Lys Tyr Lys
 465 470 475 480
 Lys Lys Phe Thr Pro Leu Trp Ser Glu Arg Tyr Ile Ser Cys Ser Arg
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 tca tta ttg gag aaa ata tct gtt gag cgt tct ttt att gaa ttt gat 96
 Ser Leu Leu Glu Lys Ile Ser Val Glu Arg Ser Phe Ile Glu Phe Asp
 20 25 30
 aaa ctt cta tta gca cct tat tgg cgt aaa gga atg ctg gca cta ata 144
 Lys Leu Leu Leu Ala Pro Tyr Trp Arg Lys Gly Met Leu Ala Leu Ile
 35 40 45
 gat agt cat gct ttt aat tat cta cca tgc tta aaa aat agg gaa tta 192
 Asp Ser His Ala Phe Asn Tyr Leu Pro Cys Leu Lys Asn Arg Glu Leu
 50 55 60
 caa tta agc gcc ttt ttg tcc cag tta gat aaa gat ttt tta ttt gag 240
 Gln Leu Ser Ala Phe Leu Ser Gln Leu Asp Lys Asp Phe Leu Phe Glu
 65 70 75 80
 aca tca gaa caa gct tgg gca tca ctc atc ttg agt atg gaa gtt gaa 288
 Thr Ser Glu Gln Ala Trp Ala Ser Leu Ile Leu Ser Met Glu Val Glu
 85 90 95

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cac aca aag act ttt tta aaa aaa tgg aag aca tca act cac ttt caa His Thr Lys Thr Phe Leu Lys Lys Trp Lys Thr Ser Thr His Phe Gln 100 105 110	336
aaa gat gtt gag cat ata gtg gat gtt tat cgt att cgt gaa caa atg Lys Asp Val Glu His Ile Val Asp Val Tyr Arg Ile Arg Glu Gln Met 115 120 125	384
gga ttg gct aaa gaa cat ctt tat cgt tat gga aaa act ata ata aaa Gly Leu Ala Lys Glu His Leu Tyr Arg Tyr Gly Lys Thr Ile Ile Lys 130 135 140	432
caa gcg gaa ggt att cgc aaa gca aga ggc ttg atg gtt gat ttc gaa Gln Ala Glu Gly Ile Arg Lys Ala Arg Gly Leu Met Val Asp Phe Glu 145 150 155 160	480
aaa ata gaa caa cta gat agt gag tta gca atc cat gat agg cat gag Lys Ile Glu Gln Leu Asp Ser Glu Leu Ala Ile His Asp Arg His Glu 165 170 175	528
ata gtt gtc aat ggt ggc acc tta atc aag aaa tta gga ata aaa cct Ile Val Val Asn Gly Gly Thr Leu Ile Lys Lys Leu Gly Ile Lys Pro 180 185 190	576
ggt cca cag atg gga gat att atc tct caa att gaa tta gcc att gtt Gly Pro Gln Met Gly Asp Ile Ile Ser Gln Ile Glu Ala Ile Val 195 200 205	624
tta gga caa ctg att aat gaa gaa gag gct att tta cat ttt gtt aag Leu Gly Gln Leu Ile Asn Glu Glu Glu Ala Ile Leu His Phe Val Lys 210 215 220	672
cag tac ttg atg gat tagagaggat tat atg agc gat ttt tta gta gat Gln Tyr Leu Met Asp Met Ser Asp Phe Leu Val Asp 225 230 235	721
gga ttg act aag tcg gtt ggt gat aag acg gtc ttt agt aat gtt tca Gly Leu Thr Lys Ser Val Gly Asp Lys Thr Val Phe Ser Asn Val Ser 240 245 250	769
ttt atc atc cat agt tta gac cgt att ggg att att ggt gtc aat gga Phe Ile Ile His Ser Leu Asp Arg Ile Gly Ile Ile Gly Val Asn Gly 255 260 265	817
act gga aag aca aca cta tta gat gtt att tcg ggt gaa tta ggt ttt Thr Gly Lys Thr Thr Leu Leu Asp Val Ile Ser Gly Glu Leu Gly Phe 270 275 280	865
gat ggt gat cgt tcc cct ttt tca tca gct aat gat tat aag att gct Asp Gly Asp Arg Ser Pro Phe Ser Ser Ala Asn Asp Tyr Lys Ile Ala 285 290 295 300	913
tat tta aaa caa gaa cca gac ttt gat gat tct cag aca att ttg gac Tyr Leu Lys Gln Glu Pro Asp Phe Asp Asp Ser Gln Thr Ile Leu Asp 305 310 315	961

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acc gta ctt tct tct gac tta aga gag atg gct tta att aaa gaa tat	1009
Thr Val Leu Ser Ser Asp Leu Arg Glu Met Ala Leu Ile Lys Glu Tyr	
320 325 330	
gaa tta ttg ctt aat cac tac gaa gaa agt aag caa tca cgt cta gag	1057
Glu Leu Leu Leu Asn His Tyr Glu Glu Ser Lys Gln Ser Arg Leu Glu	
335 340 345	
aaa gta atg gca gaa atg gat tct tta gat gct tgg tct att gag agc	1105
Lys Val Met Ala Glu Met Asp Ser Leu Asp Ala Trp Ser Ile Glu Ser	
350 355 360	
gaa gtc aaa aca gta tta tcc aaa tta ggt att act gat ttg cag ttg	1153
Glu Val Lys Thr Val Leu Ser Lys Leu Gly Ile Thr Asp Leu Gln Leu	
365 370 375 380	
tcg gtt ggt gaa tta tca gga gga tta cga aga cgt gtt caa tta gcg	1201
Ser Val Gly Glu Leu Ser Gly Gly Leu Arg Arg Arg Val Gln Leu Ala	
385 390 395	
caa gta tta tta aat gat gca gat tta ttg ctc tta gac gaa cct act	1249
Gln Val Leu Leu Asn Asp Ala Asp Leu Leu Leu Leu Asp Glu Pro Thr	
400 405 410	
aac cac tta gat att gac act att gca tgg tta acg aat ttt ttg aaa	1297
Asn His Leu Asp Ile Asp Thr Ile Ala Trp Leu Thr Asn Phe Leu Lys	
415 420 425	
aat agt aaa aag aca gtg ctt ttt ata act cat gat cgt tat ttt cta	1345
Asn Ser Lys Lys Thr Val Leu Phe Ile Thr His Asp Arg Tyr Phe Leu	
430 435 440	
gac aat gtt gca aca cgt att ttt gaa tta gat aag gca cag att aca	1393
Asp Asn Val Ala Thr Arg Ile Phe Glu Leu Asp Lys Ala Gln Ile Thr	
445 450 455 460	
gaa tat caa ggc aat tat cag gat tat gtc cga ctt cgt gca gaa caa	1441
Glu Tyr Gln Gly Asn Tyr Gln Asp Tyr Val Arg Leu Arg Ala Glu Gln	
465 470 475	
gac gag cgt gat gct gct agt tta cat aaa aag aaa cag ctt tat aaa	1489
Asp Glu Arg Asp Ala Ala Ser Leu His Lys Lys Lys Gln Leu Tyr Lys	
480 485 490	
cag gaa cta gct tgg atg cgt act cag cca caa gct cgt gca acg aaa	1537
Gln Glu Leu Ala Trp Met Arg Thr Gln Pro Gln Ala Arg Ala Thr Lys	
495 500 505	
caa cag gct cgt att aat cgt ttt caa aat cta aaa aac gat tta cac	1585
Gln Gln Ala Arg Ile Asn Arg Phe Gln Asn Leu Lys Asn Asp Leu His	
510 515 520	
caa aca agc gat aca agc gat ttg gaa atg aca ttt gaa aca agt cga	1633
Gln Thr Ser Asp Thr Ser Asp Leu Glu Met Thr Phe Glu Thr Ser Arg	
525 530 535 540	

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att ggg aaa aag gtt att aat ttt gaa aat gtc tct ttt tct tac cca Ile Gly Lys Lys Val Ile Asn Phe Glu Asn Val Ser Phe Ser Tyr Pro 545 550 555	1681
gat aaa tct atc ttg aaa gac ttt aat ttg tta att caa aat aaa gac Asp Lys Ser Ile Leu Lys Asp Phe Asn Leu Leu Ile Gln Asn Lys Asp 560 565 570	1729
cgt att ggc atc gtt gga gat aat ggt gtt gga aag tca acc tta ctt Arg Ile Gly Ile Val Gly Asp Asn Gly Val Gly Lys Ser Thr Leu Leu 575 580 585	1777
aat tta att gtt caa gat tta cag ccg gat tcg ggt aat gtc tct att Asn Leu Ile Val Gln Asp Leu Gln Pro Asp Ser Gly Asn Val Ser Ile 590 595 600	1825
ggt gaa acg ata cgt gta ggt tac ttt tca caa caa ctt cat aat atg Gly Glu Thr Ile Arg Val Gly Tyr Phe Ser Gln Gln Leu His Asn Met 605 610 615 620	1873
gat ggc tca aaa cgt gtt att aat tat ttg caa gag gtt gca gat gag Asp Gly Ser Lys Arg Val Ile Asn Tyr Leu Gln Glu Val Ala Asp Glu 625 630 635	1921
gtt aaa act agt gtc ggt aca aca agt gtg aca gaa cta ttg gaa caa Val Lys Thr Ser Val Gly Thr Thr Ser Val Thr Glu Leu Leu Glu Gln 640 645 650	1969
ttt ctc ttt cca cgt tcg aca cat gga aca caa att gca aaa tta tca Phe Leu Phe Pro Arg Ser Thr His Gly Thr Gln Ile Ala Lys Leu Ser 655 660 665	2017
ggt ggt gag aaa aaa aga ctt tac ctt tta aaa atc ctg att gaa aag Gly Gly Glu Lys Lys Arg Leu Tyr Leu Leu Lys Ile Leu Ile Glu Lys 670 675 680	2065
cct aat gtg tta cta ctt gat gag ccg aca aat gac tta gat att gct Pro Asn Val Leu Leu Leu Asp Glu Pro Thr Asn Asp Leu Asp Ile Ala 685 690 695 700	2113
aca tta act gtt ctt gaa aat ttt tta caa ggc ttt ggt ggt cct gtg Thr Leu Thr Val Leu Glu Asn Phe Leu Gln Gly Phe Gly Gly Pro Val 705 710 715	2161
att aca gtt agt cac gat cgt tac ttt tta gat aaa gtg gct aat aaa Ile Thr Val Ser His Asp Arg Tyr Phe Leu Asp Lys Val Ala Asn Lys 720 725 730	2209
att att gcg ttt gaa gat aac gat atc cgt gaa ttt ttt ggt aat tat Ile Ile Ala Phe Glu Asp Asn Asp Ile Arg Glu Phe Phe Gly Asn Tyr 735 740 745	2257
act gat tat tta gat gaa aaa gca ttt aat gag caa aat aat gaa gtt Thr Asp Tyr Leu Asp Glu Lys Ala Phe Asn Glu Gln Asn Asn Glu Val 750 755 760	2305

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atc agt aaa gag agt acc aag aca agt cgt gaa aag caa agt cgt Ile Ser Lys Lys Glu Ser Thr Lys Thr Ser Arg Glu Lys Gln Ser Arg 765 770 775 780	2353
aaa aga atg tct tac ttt gaa aaa caa gaa tgg gcg aca att gaa gac Lys Arg Met Ser Tyr Phe Glu Lys Gln Glu Trp Ala Thr Ile Glu Asp 785 790 795	2401
gat att atg ata ttg gaa aat act atc act cgt ata gaa aat gat atg Asp Ile Met Ile Leu Glu Asn Thr Ile Thr Arg Ile Glu Asn Asp Met 800 805 810	2449
caa aca tgt ggt agt gat ttt aca agg tta tct gat tta caa aag gaa Gln Thr Cys Gly Ser Asp Phe Thr Arg Leu Ser Asp Leu Gln Lys Glu 815 820 825	2497
tta gat gca aaa aat gaa gca ctt cta gaa aag tat gac cgt tat gag Leu Asp Ala Lys Asn Glu Ala Leu Leu Glu Lys Tyr Asp Arg Tyr Glu 830 835 840	2545
tac ctt agt gag ttagacac atg att atc cgt ccg att att aaa aat gat Tyr Leu Ser Glu Leu Asp Thr Met Ile Ile Arg Pro Ile Ile Lys Asn Asp 845 850 855 860	2595
gac caa gca gtt gca caa tta att cga caa agt tta cgc gcc tat gat Asp Gln Ala Val Ala Gln Leu Ile Arg Gln Ser Leu Arg Ala Tyr Asp 865 870 875	2643
tta gat aaa cct gat aca gca tat tca gac cct cac tta gat cat ttg Leu Asp Lys Pro Asp Thr Ala Tyr Ser Asp Pro His Leu Asp His Leu 880 885 890	2691
acc tca tac tac gaa aaa ata gag aag tca gga ttc ttt gtc att gag Thr Ser Tyr Tyr Glu Lys Ile Glu Lys Ser Gly Phe Phe Val Ile Glu 895 900 905	2739
gag aga gat gag att att ggc tgt ggc ggc ttt ggt ccg ctg aaa aat Glu Arg Asp Glu Ile Ile Gly Cys Gly Gly Phe Gly Pro Leu Lys Asn 910 915 920 925	2787
cta att gca gag atg cag aag gtg tac att gca gaa cgt ttc cgt ggt Leu Ile Ala Glu Met Gln Lys Val Tyr Ile Ala Glu Arg Phe Arg Gly 930 935 940	2835
aag ggg ctt gct act gat tta gtg aaa atg att gaa gta gaa gct cga Lys Gly Leu Ala Thr Asp Leu Val Lys Met Ile Glu Val Glu Ala Arg 945 950 955	2883
aaa att ggg tat aga caa ctt tat tta gag aca gcc agt act ttg agt Lys Ile Gly Tyr Arg Gln Leu Tyr Leu Glu Thr Ala Ser Thr Leu Ser 960 965 970	2931
agg gca act gcg gtt tat aag cat atg gga tat tgt gcc tta tcg caa Arg Ala Thr Ala Val Tyr Lys His Met Gly Tyr Cys Ala Leu Ser Gln 975 980 985	2979

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cca ata gca aat gat caa ggt cat aca gct atg gat att tgg atg att	3027
Pro Ile Ala Asn Asp Gln Gly His Thr Ala Met Asp Ile Trp Met Ile	
990 995 1000 1005	
aaa gat tta taagttgaaa gtggattagt gaacatggat taattatttt	3076
Lys Asp Leu	
gagataagag gaaagaaaag gagacatat atg gca tat att tgg tct tat ttg	3129
Met Ala Tyr Ile Trp Ser Tyr Leu	
1010 1015	
aaa agg tac ccc aat tgg tta tgg ctt gat tta cta gga gct atg ctt	3177
Lys Arg Tyr Pro Asn Trp Leu Trp Leu Asp Leu Leu Gly Ala Met Leu	
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ttt gtg acg gtt atc cta gga atg ccc aca gcc tta gcg ggt atg att	3225
Phe Val Thr Val Ile Leu Gly Met Pro Thr Ala Leu Ala Gly Met Ile	
1035 1040 1045	
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Asp Asn Gly Val Thr Lys Gly Asp Arg Thr Gly Val Tyr Leu Trp Thr	
1050 1055 1060	
ttc atc atg ttt ata ttt gtt gta cta ggt att att ggg cgt att acg	3321
Phe Ile Met Phe Ile Phe Val Val Leu Gly Ile Ile Gly Arg Ile Thr	
1065 1070 1075 1080	
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Met Ala Tyr Ala Ser Ser Arg Leu Thr Thr Thr Met Ile Arg Asp Met	
1085 1090 1095	
cgt aat gat atg tat gct aag ctt caa gaa tac tcc cat cat gaa tat	3417
Arg Asn Asp Met Tyr Ala Lys Leu Gln Glu Tyr Ser His His Glu Tyr	
1100 1105 1110	
gaa cag ata ggt gta tct tca cta gtg aca cgt atg aca agc gat act	3465
Glu Gln Ile Gly Val Ser Ser Leu Val Thr Arg Met Thr Ser Asp Thr	
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Phe Val Leu Met Gln Phe Ala Glu Met Ser Leu Arg Leu Gly Leu Val	
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act cct atg gta atg att ttt agc gtg gtt atg ata cta att acg agt	3561
Thr Pro Met Val Met Ile Phe Ser Val Val Met Ile Leu Ile Thr Ser	
1145 1150 1155 1160	
cca tct ttg gct tgg ctt gta gcg gtt gcg atg cct ctt ttg gta gga	3609
Pro Ser Leu Ala Trp Leu Val Ala Val Ala Met Pro Leu Leu Val Gly	
1165 1170 1175	
gtc gtt tta tat gta gct ata aaa aca aaa cct tta tct gaa aga caa	3657
Val Val Leu Tyr Val Ala Ile Lys Thr Lys Pro Leu Ser Glu Arg Gln	
1180 1185 1190	

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cag act atg ctt gat aaa atc aat caa tat gtt cgt gaa aat tta aca Gln Thr Met Leu Asp Lys Ile Asn Gln Tyr Val Arg Glu Asn Leu Thr 1195 1200 1205	3705
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gca atg att gtg gct atc gtt tgg ttt gct ttg gat ccc tta caa aga Ala Met Ile Val Ala Ile Val Trp Phe Ala Leu Asp Pro Leu Gln Arg 1260 1265 1270	3897
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cat gct ctc ttt tca ttt ttg cta ttt gcc aat ctt ttt act atg tat His Ala Leu Phe Ser Phe Leu Leu Phe Ala Asn Leu Phe Thr Met Tyr 1290 1295 1300	3993
cct cgt atg gtg gta tca agc cat cgt att aga gag gtg atg gat atg Pro Arg Met Val Val Ser Ser His Arg Ile Arg Glu Val Met Asp Met 1305 1310 1315 1320	4041
cca atc tct atc aat cct aat gcc gaa ggt gtt acg gat acg aaa ctt Pro Ile Ser Ile Asn Pro Asn Ala Glu Gly Val Thr Asp Thr Lys Leu 1325 1330 1335	4089
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aca gag agt ccc gtt ttg cat gat att tct ttt aaa gct aag cct gga Thr Glu Ser Pro Val Leu His Asp Ile Ser Phe Lys Ala Lys Pro Gly 1355 1360 1365	4185
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gtt aat ttg att cca cgt ttt tat gat gtg aca ctt gga aaa atc tta Val Asn Leu Ile Pro Arg Phe Tyr Asp Val Thr Leu Gly Lys Ile Leu 1385 1390 1395 1400	4281
gta gat gga gtt gat gta aga gat tat aac ctt aaa tca ctt cgc caa Val Asp Gly Val Asp Val Arg Asp Tyr Asn Leu Lys Ser Leu Arg Gln 1405 1410 1415	4329

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gga gag aat tta aaa tat gga aaa gct gat gct act att gat gat ctt Gly Glu Asn Leu Lys Tyr Gly Lys Ala Asp Ala Thr Ile Asp Asp Leu 1435 1440 1445	4425
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caa gaa gcc ttt gaa acg cat tta gct gaa ggt ggg agc aat ctt tct Gln Glu Ala Phe Glu Thr His Leu Ala Glu Gly Gly Ser Asn Leu Ser 1465 1470 1475 1480	4521
ggg ggt caa aaa caa cgg tta tct att gct agg gct gtt gtt aaa gat Gly Gly Gln Lys Gln Arg Leu Ser Ile Ala Arg Ala Val Val Lys Asp 1485 1490 1495	4569
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aca gac gct act tta aga gcg cgt cta aaa gaa gta acc ggt gat tct Thr Asp Ala Thr Leu Arg Ala Arg Leu Lys Glu Val Thr Gly Asp Ser 1515 1520 1525	4665
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cag att att gtc ctt gat gaa ggc gaa att gtc ggt cgt ggt acc cac Gln Ile Ile Val Leu Asp Glu Gly Glu Ile Val Gly Arg Gly Thr His 1545 1550 1555 1560	4761
gct caa tta ata gaa aat aat gct att tat cgt gaa atc gct gag tca Ala Gln Leu Ile Glu Asn Asn Ala Ile Tyr Arg Glu Ile Ala Glu Ser 1565 1570 1575	4809
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 cct tat att gct ggt att ttg att att tat ttt ttc aga ggt gtt ttc 5098
 Pro Tyr Ile Ala Gly Ile Leu Ile Ile Tyr Phe Phe Arg Gly Val Phe
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 35 40 45
 Asp Ser His Ala Phe Asn Tyr Leu Pro Cys Leu Lys Asn Arg Glu Leu
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 Gln Leu Ser Ala Phe Leu Ser Gln Leu Asp Lys Asp Phe Leu Phe Glu
 65 70 75 80
 Thr Ser Glu Gln Ala Trp Ala Ser Leu Ile Leu Ser Met Glu Val Glu
 85 90 95
 His Thr Lys Thr Phe Leu Lys Lys Trp Lys Thr Ser Thr His Phe Gln
 100 105 110
 Lys Asp Val Glu His Ile Val Asp Val Tyr Arg Ile Arg Glu Gln Met
 115 120 125
 Gly Leu Ala Lys Glu His Leu Tyr Arg Tyr Gly Lys Thr Ile Ile Lys
 130 135 140
 Gln Ala Glu Gly Ile Arg Lys Ala Arg Gly Leu Met Val Asp Phe Glu
 145 150 155 160
 Lys Ile Glu Gln Leu Asp Ser Glu Leu Ala Ile His Asp Arg His Glu
 165 170 175
 Ile Val Val Asn Gly Gly Thr Leu Ile Lys Lys Leu Gly Ile Lys Pro
 180 185 190
 Gly Pro Gln Met Gly Asp Ile Ile Ser Gln Ile Glu Leu Ala Ile Val
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 210 215 220
 Gln Tyr Leu Met Asp
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Gly	Ile	Ile	Gly	Val	Asn	Gly	Thr	Gly	Lys	Thr	Thr	Leu	Leu	Asp	Val
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Ile	Ser	Gly	Glu	Leu	Gly	Phe	Asp	Gly	Asp	Arg	Ser	Pro	Phe	Ser	Ser
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Ala	Asn	Asp	Tyr	Lys	Ile	Ala	Tyr	Leu	Lys	Gln	Glu	Pro	Asp	Phe	Asp
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Asp	Ser	Gln	Thr	Ile	Leu	Asp	Thr	Val	Leu	Ser	Ser	Asp	Leu	Arg	Glu
				85					90					95	
Met	Ala	Leu	Ile	Lys	Glu	Tyr	Glu	Leu	Leu	Leu	Asn	His	Tyr	Glu	Glu
			100					105					110		
Ser	Lys	Gln	Ser	Arg	Leu	Glu	Lys	Val	Met	Ala	Glu	Met	Asp	Ser	Leu
		115					120					125			
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Gly	Ile	Thr	Asp	Leu	Gln	Leu	Ser	Val	Gly	Glu	Leu	Ser	Gly	Gly	Leu
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Arg	Arg	Arg	Val	Gln	Leu	Ala	Gln	Val	Leu	Leu	Asn	Asp	Ala	Asp	Leu
				165					170					175	
Leu	Leu	Leu	Asp	Glu	Pro	Thr	Asn	His	Leu	Asp	Ile	Asp	Thr	Ile	Ala
			180					185					190		
Trp	Leu	Thr	Asn	Phe	Leu	Lys	Asn	Ser	Lys	Lys	Thr	Val	Leu	Phe	Ile
		195					200					205			
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Leu	Asp	Lys	Ala	Gln	Ile	Thr	Glu	Tyr	Gln	Gly	Asn	Tyr	Gln	Asp	Tyr
225					230					235					240
Val	Arg	Leu	Arg	Ala	Glu	Gln	Asp	Glu	Arg	Asp	Ala	Ala	Ser	Leu	His
			245						250					255	
Lys	Lys	Lys	Gln	Leu	Tyr	Lys	Gln	Glu	Leu	Ala	Trp	Met	Arg	Thr	Gln
			260					265					270		
Pro	Gln	Ala	Arg	Ala	Thr	Lys	Gln	Gln	Ala	Arg	Ile	Asn	Arg	Phe	Gln
		275					280					285			
Asn	Leu	Lys	Asn	Asp	Leu	His	Gln	Thr	Ser	Asp	Thr	Ser	Asp	Leu	Glu
	290					295					300				
Met	Thr	Phe	Glu	Thr	Ser	Arg	Ile	Gly	Lys	Lys	Val	Ile	Asn	Phe	Glu
305					310					315					320
Asn	Val	Ser	Phe	Ser	Tyr	Pro	Asp	Lys	Ser	Ile	Leu	Lys	Asp	Phe	Asn
			325						330					335	
Leu	Leu	Ile	Gln	Asn	Lys	Asp	Arg	Ile	Gly	Ile	Val	Gly	Asp	Asn	Gly
		340						345					350		
Val	Gly	Lys	Ser	Thr	Leu	Leu	Asn	Leu	Ile	Val	Gln	Asp	Leu	Gln	Pro
		355					360					365			
Asp	Ser	Gly	Asn	Val	Ser	Ile	Gly	Glu	Thr	Ile	Arg	Val	Gly	Tyr	Phe
	370					375					380				
Ser	Gln	Gln	Leu	His	Asn	Met	Asp	Gly	Ser	Lys	Arg	Val	Ile	Asn	Tyr
385					390					395					400
Leu	Gln	Glu	Val	Ala	Asp	Glu	Val	Lys	Thr	Ser	Val	Gly	Thr	Thr	Ser
			405						410					415	
Val	Thr	Glu	Leu	Leu	Glu	Gln	Phe	Leu	Phe	Pro	Arg	Ser	Thr	His	Gly
			420					425						430	

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Thr Gln Ile Ala Lys Leu Ser Gly Gly Glu Lys Lys Arg Leu Tyr Leu
 435 440 445
 Leu Lys Ile Leu Ile Glu Lys Pro Asn Val Leu Leu Asp Glu Pro
 450 455 460
 Thr Asn Asp Leu Asp Ile Ala Thr Leu Thr Val Leu Glu Asn Phe Leu
 465 470 475 480
 Gln Gly Phe Gly Gly Pro Val Ile Thr Val Ser His Asp Arg Tyr Phe
 485 490 495
 Leu Asp Lys Val Ala Asn Lys Ile Ile Ala Phe Glu Asp Asn Asp Ile
 500 505 510
 Arg Glu Phe Phe Gly Asn Tyr Thr Asp Tyr Leu Asp Glu Lys Ala Phe
 515 520 525
 Asn Glu Gln Asn Asn Glu Val Ile Ser Lys Lys Glu Ser Thr Lys Thr
 530 535 540
 Ser Arg Glu Lys Gln Ser Arg Lys Arg Met Ser Tyr Phe Glu Lys Gln
 545 550 555 560
 Glu Trp Ala Thr Ile Glu Asp Asp Ile Met Ile Leu Glu Asn Thr Ile
 565 570 575
 Thr Arg Ile Glu Asn Asp Met Gln Thr Cys Gly Ser Asp Phe Thr Arg
 580 585 590
 Leu Ser Asp Leu Gln Lys Glu Leu Asp Ala Lys Asn Glu Ala Leu Leu
 595 600 605
 Glu Lys Tyr Asp Arg Tyr Glu Tyr Leu Ser Glu Leu Asp Thr
 610 615 620

<210> 10
 <211> 157
 <212> PRT
 <213> Streptococcus

<400> 10
 Met Ile Ile Arg Pro Ile Ile Lys Asn Asp Asp Gln Ala Val Ala Gln
 1 5 10 15
 Leu Ile Arg Gln Ser Leu Arg Ala Tyr Asp Leu Asp Lys Pro Asp Thr
 20 25 30
 Ala Tyr Ser Asp Pro His Leu Asp His Leu Thr Ser Tyr Tyr Glu Lys
 35 40 45
 Ile Glu Lys Ser Gly Phe Phe Val Ile Glu Glu Arg Asp Glu Ile Ile
 50 55 60
 Gly Cys Gly Gly Phe Gly Pro Leu Lys Asn Leu Ile Ala Glu Met Gln
 65 70 75 80
 Lys Val Tyr Ile Ala Glu Arg Phe Arg Gly Lys Gly Leu Ala Thr Asp
 85 90 95
 Leu Val Lys Met Ile Glu Val Glu Ala Arg Lys Ile Gly Tyr Arg Gln
 100 105 110
 Leu Tyr Leu Glu Thr Ala Ser Thr Leu Ser Arg Ala Thr Ala Val Tyr
 115 120 125
 Lys His Met Gly Tyr Cys Ala Leu Ser Gln Pro Ile Ala Asn Asp Gln
 130 135 140
 Gly His Thr Ala Met Asp Ile Trp Met Ile Lys Asp Leu
 145 150 155

<210> 11
 <211> 579
 <212> PRT
 <213> Streptococcus

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<400> 11

Met	Ala	Tyr	Ile	Trp	Ser	Tyr	Leu	Lys	Arg	Tyr	Pro	Asn	Trp	Leu	Trp
1				5					10					15	
Leu	Asp	Leu	Leu	Gly	Ala	Met	Leu	Phe	Val	Thr	Val	Ile	Leu	Gly	Met
			20					25					30		
Pro	Thr	Ala	Leu	Ala	Gly	Met	Ile	Asp	Asn	Gly	Val	Thr	Lys	Gly	Asp
		35					40					45			
Arg	Thr	Gly	Val	Tyr	Leu	Trp	Thr	Phe	Ile	Met	Phe	Ile	Phe	Val	Val
	50					55					60				
Leu	Gly	Ile	Ile	Gly	Arg	Ile	Thr	Met	Ala	Tyr	Ala	Ser	Ser	Arg	Leu
65					70					75				80	
Thr	Thr	Thr	Met	Ile	Arg	Asp	Met	Arg	Asn	Asp	Met	Tyr	Ala	Lys	Leu
				85					90					95	
Gln	Glu	Tyr	Ser	His	His	Glu	Tyr	Glu	Gln	Ile	Gly	Val	Ser	Ser	Leu
			100					105					110		
Val	Thr	Arg	Met	Thr	Ser	Asp	Thr	Phe	Val	Leu	Met	Gln	Phe	Ala	Glu
		115					120					125			
Met	Ser	Leu	Arg	Leu	Gly	Leu	Val	Thr	Pro	Met	Val	Met	Ile	Phe	Ser
	130					135					140				
Val	Val	Met	Ile	Leu	Ile	Thr	Ser	Pro	Ser	Leu	Ala	Trp	Leu	Val	Ala
145					150					155					160
Val	Ala	Met	Pro	Leu	Leu	Val	Gly	Val	Val	Leu	Tyr	Val	Ala	Ile	Lys
				165					170					175	
Thr	Lys	Pro	Leu	Ser	Glu	Arg	Gln	Gln	Thr	Met	Leu	Asp	Lys	Ile	Asn
			180					185					190		
Gln	Tyr	Val	Arg	Glu	Asn	Leu	Thr	Gly	Leu	Arg	Val	Val	Arg	Ala	Phe
		195					200					205			
Ala	Arg	Glu	Asn	Phe	Gln	Ser	Gln	Lys	Phe	Gln	Val	Ala	Asn	Gln	Arg
	210					215					220				
Tyr	Thr	Asp	Thr	Ser	Thr	Gly	Leu	Phe	Lys	Leu	Thr	Gly	Leu	Thr	Glu
225					230					235					240
Pro	Leu	Phe	Val	Gln	Ile	Ile	Ile	Ala	Met	Ile	Val	Ala	Ile	Val	Trp
				245					250				255		
Phe	Ala	Leu	Asp	Pro	Leu	Gln	Arg	Gly	Ala	Ile	Lys	Ile	Gly	Asp	Leu
			260					265					270		
Val	Ala	Phe	Ile	Glu	Tyr	Ser	Phe	His	Ala	Leu	Phe	Ser	Phe	Leu	Leu
		275					280					285			
Phe	Ala	Asn	Leu	Phe	Thr	Met	Tyr	Pro	Arg	Met	Val	Val	Ser	Ser	His
	290					295					300				
Arg	Ile	Arg	Glu	Val	Met	Asp	Met	Pro	Ile	Ser	Ile	Asn	Pro	Asn	Ala
305					310					315					320
Glu	Gly	Val	Thr	Asp	Thr	Lys	Leu	Lys	Gly	His	Leu	Glu	Phe	Asp	Asn
				325					330					335	
Val	Thr	Phe	Ala	Tyr	Pro	Gly	Glu	Thr	Glu	Ser	Pro	Val	Leu	His	Asp
			340					345					350		
Ile	Ser	Phe	Lys	Ala	Lys	Pro	Gly	Glu	Thr	Ile	Ala	Phe	Ile	Gly	Ser
		355					360					365			
Thr	Gly	Ser	Gly	Lys	Ser	Ser	Leu	Val	Asn	Leu	Ile	Pro	Arg	Phe	Tyr
	370					375					380				
Asp	Val	Thr	Leu	Gly	Lys	Ile	Leu	Val	Asp	Gly	Val	Asp	Val	Arg	Asp
385					390					395					400
Tyr	Asn	Leu	Lys	Ser	Leu	Arg	Gln	Lys	Ile	Gly	Phe	Ile	Pro	Gln	Lys
				405						410				415	
Ala	Leu	Leu	Phe	Thr	Gly	Thr	Ile	Gly	Glu	Asn	Leu	Lys	Tyr	Gly	Lys
			420					425						430	

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Ala Asp Ala Thr Ile Asp Asp Leu Arg Gln Ala Val Asp Ile Ser Gln
 435 440 445
 Ala Lys Glu Phe Ile Glu Ser His Gln Glu Ala Phe Glu Thr His Leu
 450 455 460
 Ala Glu Gly Gly Ser Asn Leu Ser Gly Gly Gln Lys Gln Arg Leu Ser
 465 470 475 480
 Ile Ala Arg Ala Val Val Lys Asp Pro Asp Leu Tyr Ile Phe Asp Asp
 485 490 495
 Ser Phe Ser Ala Leu Asp Tyr Lys Thr Asp Ala Thr Leu Arg Ala Arg
 500 505 510
 Leu Lys Glu Val Thr Gly Asp Ser Thr Val Leu Ile Val Ala Gln Arg
 515 520 525
 Val Gly Thr Ile Met Asp Ala Asp Gln Ile Ile Val Leu Asp Glu Gly
 530 535 540
 Glu Ile Val Gly Arg Gly Thr His Ala Gln Leu Ile Glu Asn Asn Ala
 545 550 555 560
 Ile Tyr Arg Glu Ile Ala Glu Ser Gln Leu Lys Asn Gln Asn Leu Ser
 565 570 575
 Glu Gly Glu

<210> 12
 <211> 92
 <212> PRT
 <213> Streptococcus

<400> 12
 Met Arg Lys Lys Ser Val Phe Leu Arg Leu Trp Ser Tyr Leu Thr Arg
 1 5 10 15
 Tyr Lys Ala Thr Leu Phe Leu Ala Ile Phe Leu Lys Val Leu Ser Ser
 20 25 30
 Phe Met Ser Val Leu Glu Pro Phe Ile Leu Gly Leu Ala Ile Thr Glu
 35 40 45
 Leu Thr Ala Asn Leu Val Asp Met Ala Lys Gly Val Ser Gly Ala Glu
 50 55 60
 Leu Asn Val Pro Tyr Ile Ala Gly Ile Leu Ile Ile Tyr Phe Phe Arg
 65 70 75 80
 Gly Val Phe Tyr Glu Leu Gly Ser Tyr Gly Ser Asn
 85 90

<210> 13
 <211> 5215
 <212> DNA
 <213> Streptococcus

<220>
 <221> CDS
 <222> (3)...(122)
 <221> CDS
 <222> (133)...(2511)
 <221> CDS
 <222> (367)...(2511)
 <221> CDS

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<222> (2946)...(2716)
 <223> of complementary strand

<221> CDS
 <222> (3252)...(2995)
 <223> of complementary strand

<221> CDS
 <222> (3676)...(3299)
 <223> of complementary strand

<221> CDS
 <222> (4124)...(3837)
 <223> of complementary strand

<221> CDS
 <222> (5214)...(4351)
 <223> of complementary strand

<400> 13

aa ttt gga agt gct cta tca aca gtt gaa gta aag gag att att agt	47
Phe Gly Ser Ala Leu Ser Thr Val Glu Val Lys Glu Ile Ile Ser	
1 5 10 15	
 gaa gaa aac ata tgg tta tat cgg ctc agt tgc tgc cat ttt act agc	95
Glu Glu Asn Ile Trp Leu Tyr Arg Leu Ser Cys Cys His Phe Thr Ser	
20 25 30	
 tac tca tat tgg aag tta cca act tgg taagcatcat atg ggt cta gca	144
Tyr Ser Tyr Trp Lys Leu Pro Thr Trp Met Gly Leu Ala	
35 40	
 aca aag gac aat cag att gcc tat att gat gac agc aaa ggt aag gca	192
Thr Lys Asp Asn Gln Ile Ala Tyr Ile Asp Asp Ser Lys Gly Lys Ala	
45 50 55 60	
 aaa gcc cct aaa aca aac aaa acg atg gat caa atc agt gct gaa gaa	240
Lys Ala Pro Lys Thr Asn Lys Thr Met Asp Gln Ile Ser Ala Glu Glu	
65 70 75	
 ggc atc tct gct gaa cag atc gta gtc aaa att act gac caa ggc tat	288
Gly Ile Ser Ala Glu Gln Ile Val Val Lys Ile Thr Asp Gln Gly Tyr	
80 85 90	
 gtg acc tca cac ggt gac cat tat cat ttt tac aat ggg aaa gtt cct	336
Val Thr Ser His Gly Asp His Tyr His Phe Tyr Asn Gly Lys Val Pro	
95 100 105	
 tat gat gcg att att agt gaa gag ttg ttg atg acg gat cct aat tac	384
Tyr Asp Ala Ile Ile Ser Glu Glu Leu Leu Met Thr Asp Pro Asn Tyr	
110 115 120	
 cgt ttt aaa caa tca gac gtt atc aat gaa atc tta gac ggt tac gtt	432
Arg Phe Lys Gln Ser Asp Val Ile Asn Glu Ile Leu Asp Gly Tyr Val	
125 130 135 140	

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att aaa gtc aat ggc aac tat tat gtt tac ctc aag cca ggt agt aag Ile Lys Val Asn Gly Asn Tyr Tyr Val Tyr Leu Lys Pro Gly Ser Lys 145 150 155	480
cgc aaa aac att cga acc aaa caa caa att gct gag caa gta gcc aaa Arg Lys Asn Ile Arg Thr Lys Gln Gln Ile Ala Glu Gln Val Ala Lys 160 165 170	528
gga act aaa gaa gct aaa gaa aaa ggt tta gct caa gtg gcc cat ctc Gly Thr Lys Glu Ala Lys Glu Lys Gly Leu Ala Gln Val Ala His Leu 175 180 185	576
agt aaa gaa gaa gtt gcg gca gtc aat gaa gca aaa aga caa gga cgc Ser Lys Glu Glu Val Ala Ala Val Asn Glu Ala Lys Arg Gln Gly Arg 190 195 200	624
tat act aca gac gat ggc tat att ttt agt ccg aca gat atc att gat Tyr Thr Thr Asp Asp Gly Tyr Ile Phe Ser Pro Thr Asp Ile Ile Asp 205 210 215 220	672
gat tta gga gat gct tat tta gta cct cat ggt aat cac tat cat tat Asp Leu Gly Asp Ala Tyr Leu Val Pro His Gly Asn His Tyr His Tyr 225 230 235	720
att cct aaa aag gat ttg tct cca agt gag cta gct gct gca caa gcc Ile Pro Lys Lys Asp Leu Ser Pro Ser Glu Leu Ala Ala Ala Gln Ala 240 245 250	768
tac tgg agt caa aaa caa ggt cga ggt gct aga ccg tct gat tac cgc Tyr Trp Ser Gln Lys Gln Gly Arg Gly Ala Arg Pro Ser Asp Tyr Arg 255 260 265	816
ccg aca cca gcc cca ggt cgt agg aaa gcc cca att cct gat gtg acg Pro Thr Pro Ala Pro Gly Arg Arg Lys Ala Pro Ile Pro Asp Val Thr 270 275 280	864
cct aac cct gga caa ggt cat cag cca gat aac ggt ggc tat cat cca Pro Asn Pro Gly Gln Gly His Gln Pro Asp Asn Gly Gly Tyr His Pro 285 290 295 300	912
gcg cct cct agg cca aat gat gcg tca caa aac aaa cac caa aga gat Ala Pro Pro Arg Pro Asn Asp Ala Ser Gln Asn Lys His Gln Arg Asp 305 310 315	960
gag ttt aaa gga aaa acc ttt aag gaa ctt tta gat caa cta cac cgt Glu Phe Lys Gly Lys Thr Phe Lys Glu Leu Leu Asp Gln Leu His Arg 320 325 330	1008
ctt gat ttg aaa tac cgt cat gtg gaa gaa gat ggg ttg att ttt gaa Leu Asp Leu Lys Tyr Arg His Val Glu Glu Asp Gly Leu Ile Phe Glu 335 340 345	1056
ccg act caa gtg atc aaa tca aac gct ttt ggg tat gtg gtg cct cat Pro Thr Gln Val Ile Lys Ser Asn Ala Phe Gly Tyr Val Val Pro His 350 355 360	1104

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gga gat cat tat cat att atc cca aga agt cag tta tca cct ctt gaa	1152
Gly Asp His Tyr His Ile Ile Pro Arg Ser Gln Leu Ser Pro Leu Glu	
365 370 375 380	
atg gaa tta gca gat cga tac tta gct ggc caa act gag gac aat gac	1200
Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly Gln Thr Glu Asp Asn Asp	
385 390 395	
tca ggt tca gag cac tca aaa cca tca gat aaa gaa gtg aca cat acc	1248
Ser Gly Ser Glu His Ser Lys Pro Ser Asp Lys Glu Val Thr His Thr	
400 405 410	
ttt ctt ggt cat cgc atc aaa gct tac gga aaa ggc tta gat ggt aaa	1296
Phe Leu Gly His Arg Ile Lys Ala Tyr Gly Lys Gly Leu Asp Gly Lys	
415 420 425	
cca tat gat acg agt gat gct tat gtt ttt agt aaa gaa tcc att cat	1344
Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe Ser Lys Glu Ser Ile His	
430 435 440	
tca gtg gat aaa tca gga gtt aca gct aaa cac gga gat cat ttc cac	1392
Ser Val Asp Lys Ser Gly Val Thr Ala Lys His Gly Asp His Phe His	
445 450 455 460	
tat ata gga ttt gga gaa ctt gaa caa tat gag ttg gat gag gtc gct	1440
Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr Glu Leu Asp Glu Val Ala	
465 470 475	
aac tgg gtg aaa gca aaa ggt caa gct gat gag ctt gct gct gct ttg	1488
Asn Trp Val Lys Ala Lys Gly Gln Ala Asp Glu Leu Ala Ala Ala Leu	
480 485 490	
gat cag gaa caa ggc aaa gaa aaa cca ctc ttt gac act aaa aaa gtg	1536
Asp Gln Glu Gln Gly Lys Glu Lys Pro Leu Phe Asp Thr Lys Lys Val	
495 500 505	
agt cgc aaa gta aca aaa gat ggt aaa gtg ggc tat atg atg cca aaa	1584
Ser Arg Lys Val Thr Lys Asp Gly Lys Val Gly Tyr Met Met Pro Lys	
510 515 520	
gat ggt aag gac tat ttc tat gct cgt gat caa ctt gat ttg act cag	1632
Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp Gln Leu Asp Leu Thr Gln	
525 530 535 540	
att gcc ttt gcc gaa caa gaa cta atg ctt aaa gat aag aag cat tac	1680
Ile Ala Phe Ala Glu Gln Glu Leu Met Leu Lys Asp Lys Lys His Tyr	
545 550 555	
cgt tat gac att gtt gac aca ggt att gag cca cga ctt gct gta gat	1728
Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu Pro Arg Leu Ala Val Asp	
560 565 570	
gtg tca agt ctg ccg atg cat gct ggt aat gct act tac gat act gga	1776
Val Ser Ser Leu Pro Met His Ala Gly Asn Ala Thr Tyr Asp Thr Gly	
575 580 585	

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agt tcg ttt gtt atc cca cat att gat cat atc cat gtc gtt ccg tat	1824
Ser Ser Phe Val Ile Pro His Ile Asp His Ile His Val Val Pro Tyr	
590 595 600	
tca tgg ttg acg cgc gat cag att gca aca gtc aag tat gtg atg caa	1872
Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr Val Lys Tyr Val Met Gln	
605 610 615 620	
cac ccc gaa gtt cgt ccg gat gta tgg tct aag cca ggg cat gaa gag	1920
His Pro Glu Val Arg Pro Asp Val Trp Ser Lys Pro Gly His Glu Glu	
625 630 635	
tca ggt tcg gtc att cca aat gtt acg cct ctt gat aaa cgt gct ggt	1968
Ser Gly Ser Val Ile Pro Asn Val Thr Pro Leu Asp Lys Arg Ala Gly	
640 645 650	
atg cca aac tgg caa att atc cat tct gct gaa gaa gtt caa aaa gcc	2016
Met Pro Asn Trp Gln Ile Ile His Ser Ala Glu Glu Val Gln Lys Ala	
655 660 665	
cta gca gaa ggt cgt ttt gca aca cca gac ggc tat att ttc gat cca	2064
Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp Gly Tyr Ile Phe Asp Pro	
670 675 680	
cga gat gtt ttg gcc aaa gaa act ttt gta tgg aaa gat ggc tcc ttt	2112
Arg Asp Val Leu Ala Lys Glu Thr Phe Val Trp Lys Asp Gly Ser Phe	
685 690 695 700	
agc atc cca aga gca gat ggc agt tca ttg aga acc att aat aaa tct	2160
Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu Arg Thr Ile Asn Lys Ser	
705 710 715	
gat cta tcc caa gct gag tgg caa caa gct caa gag tta ttg gca aag	2208
Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala Gln Glu Leu Leu Ala Lys	
720 725 730	
aaa aat act ggt gat gct act gat acg gat aaa ccc aaa gaa aag caa	2256
Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp Lys Pro Lys Glu Lys Gln	
735 740 745	
cag gca gat aag agc aat gaa aac caa cag cca agt gaa gcc agt aaa	2304
Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln Pro Ser Glu Ala Ser Lys	
750 755 760	
gaa gaa aaa gaa tca gat gac ttt ata gac agt tta cca gac tat ggt	2352
Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp Ser Leu Pro Asp Tyr Gly	
765 770 775 780	
cta gat aga gca acc cta gaa gat cat atc aat caa tta gca caa aaa	2400
Leu Asp Arg Ala Thr Leu Glu Asp His Ile Asn Gln Leu Ala Gln Lys	
785 790 795	
gct aat atc gat cct aag tat ctc att ttc caa cca gaa ggt gtc caa	2448
Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe Gln Pro Glu Gly Val Gln	
800 805 810	

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ttt tat aat aaa aat ggt gaa ttg gta act tat gat atc aag aca ctt 2496
 Phe Tyr Asn Lys Asn Gly Glu Leu Val Thr Tyr Asp Ile Lys Thr Leu
 815 820 825

caa caa ata aac cct taacccaaaag aagatctcat tgttaaagca ctgctttgtc 2551
 Gln Gln Ile Asn Pro
 830

aaagcaagtt acggtgattt tgaagtcatt ctatgtaacg agtagtgata aaagttggat 2611
 aatagcgggt ttcttttgca aagaaatggg atccatgtta gaatagtaaa aaaagaggag 2671
 gattcttggg ctaatgtcaa ataagtagac agaaaactgt gttattttattgcgt 2726
 taaaataatt ttcttctttc tgattagggg ttagtcctag attagccgta tgtgggttgt 2786
 aattgttata aaaattctca atgtattcaa agcagtcctaa ttgaacctgt ttgatatttt 2846
 gataatgttt tcggttgatt tgtctatgct ttaaatactt gaaaaatgct tcagttacgg 2906
 cattatcata aggatatcca ggattagaaa aagaatgcat gatattggca ctgcacccta 2966
 atagtgaagc gcaagaaaaa cacttttaggcaatcagtt ttctgtactg tacaggcgac 3025
 tggtcgttta atctctggtt aattctagtt tcattataaa atgtaatgta atttttaaca 3085
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 gagccttggt cactgtgtaa gattgtcctt ttatttaggcaatt ttaactgatt 3319
 aaggggtgct agtacaaaat ccgtgtcctg acaatctgag atagtgtgag ctataatttc 3379
 tcggttatag agattcataa ttgatgagag atacaattta cagttaccga aatataggta 3439
 ggtaatatct gttacgagct ttctcttagg cttatcggca tggaaatccc gactcaattt 3499
 attatctggt aaataataag ctttacccaa attgggaact ttcttggtac gtgtccgaca 3559
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 gatttttttg agcaatcgtg taatggtacg atagccataa ataaagtgat tctccatata 3679
 gagctgttca attaatcaa taaggctatc tttttttgcg gcttctcata ctcttttttc 3739
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 cggctcttggt ccaaccaaga gacaggccga tgatttcacg gttgtatagg tcaatgatga 4751
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 tcataatgag tcgaatccga cgacgtgaaa gtgtgatacc ttcgttattc aagcatattt 4991
 tgatttttct ggatccgtat ctgactcgc tatcgagaaa aattctttta atagtttctt 5051
 caaactccgt ttcagatact gactccacgg cttgatagta ataacttgag tgtggcatat 5111
 tcagccagcg acacatcttt gaaatgctgt atttatcctt attagcagtg attatttccc 5171
 tttttgtgcc ataaccacg ctgcttgctt taggatatct aatt 5215

<210> 14

<211> 40

<212> PRT

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<213> Streptococcus

<400> 14

Phe Gly Ser Ala Leu Ser Thr Val Glu Val Lys Glu Ile Ile Ser Glu
 1 5 10 15
 Glu Asn Ile Trp Leu Tyr Arg Leu Ser Cys Cys His Phe Thr Ser Tyr
 20 25 30
 Ser Tyr Trp Lys Leu Pro Thr Trp
 35 40

<210> 15

<211> 793

<212> PRT

<213> Streptococcus

<400> 15

Met Gly Leu Ala Thr Lys Asp Asn Gln Ile Ala Tyr Ile Asp Asp Ser
 1 5 10 15
 Lys Gly Lys Ala Lys Ala Pro Lys Thr Asn Lys Thr Met Asp Gln Ile
 20 25 30
 Ser Ala Glu Glu Gly Ile Ser Ala Glu Gln Ile Val Val Lys Ile Thr
 35 40 45
 Asp Gln Gly Tyr Val Thr Ser His Gly Asp His Tyr His Phe Tyr Asn
 50 55 60
 Gly Lys Val Pro Tyr Asp Ala Ile Ile Ser Glu Glu Leu Leu Met Thr
 65 70 75 80
 Asp Pro Asn Tyr Arg Phe Lys Gln Ser Asp Val Ile Asn Glu Ile Leu
 85 90 95
 Asp Gly Tyr Val Ile Lys Val Asn Gly Asn Tyr Tyr Val Tyr Leu Lys
 100 105 110
 Pro Gly Ser Lys Arg Lys Asn Ile Arg Thr Lys Gln Gln Ile Ala Glu
 115 120 125
 Gln Val Ala Lys Gly Thr Lys Glu Ala Lys Glu Lys Gly Leu Ala Gln
 130 135 140
 Val Ala His Leu Ser Lys Glu Glu Val Ala Ala Val Asn Glu Ala Lys
 145 150 155 160
 Arg Gln Gly Arg Tyr Thr Thr Asp Asp Gly Tyr Ile Phe Ser Pro Thr
 165 170 175
 Asp Ile Ile Asp Asp Leu Gly Asp Ala Tyr Leu Val Pro His Gly Asn
 180 185 190
 His Tyr His Tyr Ile Pro Lys Lys Asp Leu Ser Pro Ser Glu Leu Ala
 195 200 205
 Ala Ala Gln Ala Tyr Trp Ser Gln Lys Gln Gly Arg Gly Ala Arg Pro
 210 215 220
 Ser Asp Tyr Arg Pro Thr Pro Ala Pro Gly Arg Arg Lys Ala Pro Ile
 225 230 235 240
 Pro Asp Val Thr Pro Asn Pro Gly Gln Gly His Gln Pro Asp Asn Gly
 245 250 255
 Gly Tyr His Pro Ala Pro Pro Arg Pro Asn Asp Ala Ser Gln Asn Lys
 260 265 270
 His Gln Arg Asp Glu Phe Lys Gly Lys Thr Phe Lys Glu Leu Leu Asp
 275 280 285
 Gln Leu His Arg Leu Asp Leu Lys Tyr Arg His Val Glu Glu Asp Gly
 290 295 300
 Leu Ile Phe Glu Pro Thr Gln Val Ile Lys Ser Asn Ala Phe Gly Tyr
 305 310 315 320

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Val Val Pro His Gly Asp His Tyr His Ile Ile Pro Arg Ser Gln Leu
 325 330 335
 Ser Pro Leu Glu Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly Gln Thr
 340 345 350
 Glu Asp Asn Asp Ser Gly Ser Glu His Ser Lys Pro Ser Asp Lys Glu
 355 360 365
 Val Thr His Thr Phe Leu Gly His Arg Ile Lys Ala Tyr Gly Lys Gly
 370 375 380
 Leu Asp Gly Lys Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe Ser Lys
 385 390 395 400
 Glu Ser Ile His Ser Val Asp Lys Ser Gly Val Thr Ala Lys His Gly
 405 410 415
 Asp His Phe His Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr Glu Leu
 420 425 430
 Asp Glu Val Ala Asn Trp Val Lys Ala Lys Gly Gln Ala Asp Glu Leu
 435 440 445
 Ala Ala Ala Leu Asp Gln Glu Gly Lys Glu Lys Pro Leu Phe Asp
 450 455 460
 Thr Lys Lys Val Ser Arg Lys Val Thr Lys Asp Gly Lys Val Gly Tyr
 465 470 475 480
 Met Met Pro Lys Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp Gln Leu
 485 490 495
 Asp Leu Thr Gln Ile Ala Phe Ala Glu Gln Glu Leu Met Leu Lys Asp
 500 505 510
 Lys Lys His Tyr Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu Pro Arg
 515 520 525
 Leu Ala Val Asp Val Ser Ser Leu Pro Met His Ala Gly Asn Ala Thr
 530 535 540
 Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His Ile His
 545 550 555 560
 Val Val Pro Tyr Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr Val Lys
 565 570 575
 Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser Lys Pro
 580 585 590
 Gly His Glu Glu Ser Gly Ser Val Ile Pro Asn Val Thr Pro Leu Asp
 595 600 605
 Lys Arg Ala Gly Met Pro Asn Trp Gln Ile Ile His Ser Ala Glu Glu
 610 615 620
 Val Gln Lys Ala Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp Gly Tyr
 625 630 635 640
 Ile Phe Asp Pro Arg Asp Val Leu Ala Lys Glu Thr Phe Val Trp Lys
 645 650 655
 Asp Gly Ser Phe Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu Arg Thr
 660 665 670
 Ile Asn Lys Ser Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala Gln Glu
 675 680 685
 Leu Leu Ala Lys Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp Lys Pro
 690 695 700
 Lys Glu Lys Gln Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln Pro Ser
 705 710 715 720
 Glu Ala Ser Lys Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp Ser Leu
 725 730 735
 Pro Asp Tyr Gly Leu Asp Arg Ala Thr Leu Glu Asp His Ile Asn Gln
 740 745 750
 Leu Ala Gln Lys Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe Gln Pro
 755 760 765

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Glu Gly Val Gln Phe Tyr Asn Lys Asn Gly Glu Leu Val Thr Tyr Asp
 770 775 780
 Ile Lys Thr Leu Gln Gln Ile Asn Pro
 785 790

<210> 16
 <211> 715
 <212> PRT
 <213> Streptococcus

<400> 16
 Met Thr Asp Pro Asn Tyr Arg Phe Lys Gln Ser Asp Val Ile Asn Glu
 1 5 10 15
 Ile Leu Asp Gly Tyr Val Ile Lys Val Asn Gly Asn Tyr Tyr Val Tyr
 20 25 30
 Leu Lys Pro Gly Ser Lys Arg Lys Asn Ile Arg Thr Lys Gln Gln Ile
 35 40 45
 Ala Glu Gln Val Ala Lys Gly Thr Lys Glu Ala Lys Glu Lys Gly Leu
 50 55 60
 Ala Gln Val Ala His Leu Ser Lys Glu Glu Val Ala Ala Val Asn Glu
 65 70 75 80
 Ala Lys Arg Gln Gly Arg Tyr Thr Thr Asp Asp Gly Tyr Ile Phe Ser
 85 90 95
 Pro Thr Asp Ile Ile Asp Asp Leu Gly Asp Ala Tyr Leu Val Pro His
 100 105 110
 Gly Asn His Tyr His Tyr Ile Pro Lys Lys Asp Leu Ser Pro Ser Glu
 115 120 125
 Leu Ala Ala Ala Gln Ala Tyr Trp Ser Gln Lys Gln Gly Arg Gly Ala
 130 135 140
 Arg Pro Ser Asp Tyr Arg Pro Thr Pro Ala Pro Gly Arg Arg Lys Ala
 145 150 155 160
 Pro Ile Pro Asp Val Thr Pro Asn Pro Gly Gln Gly His Gln Pro Asp
 165 170 175
 Asn Gly Gly Tyr His Pro Ala Pro Pro Arg Pro Asn Asp Ala Ser Gln
 180 185 190
 Asn Lys His Gln Arg Asp Glu Phe Lys Gly Lys Thr Phe Lys Glu Leu
 195 200 205
 Leu Asp Gln Leu His Arg Leu Asp Leu Lys Tyr Arg His Val Glu Glu
 210 215 220
 Asp Gly Leu Ile Phe Glu Pro Thr Gln Val Ile Lys Ser Asn Ala Phe
 225 230 235 240
 Gly Tyr Val Val Pro His Gly Asp His Tyr His Ile Ile Pro Arg Ser
 245 250 255
 Gln Leu Ser Pro Leu Glu Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly
 260 265 270
 Gln Thr Glu Asp Asn Asp Ser Gly Ser Glu His Ser Lys Pro Ser Asp
 275 280 285
 Lys Glu Val Thr His Thr Phe Leu Gly His Arg Ile Lys Ala Tyr Gly
 290 295 300
 Lys Gly Leu Asp Gly Lys Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe
 305 310 315 320
 Ser Lys Glu Ser Ile His Ser Val Asp Lys Ser Gly Val Thr Ala Lys
 325 330 335
 His Gly Asp His Phe His Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr
 340 345 350

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Glu Leu Asp Glu Val Ala Asn Trp Val Lys Ala Lys Gly Gln Ala Asp
 355 360 365
 Glu Leu Ala Ala Ala Leu Asp Gln Glu Gln Gly Lys Glu Lys Pro Leu
 370 375 380
 Phe Asp Thr Lys Lys Val Ser Arg Lys Val Thr Lys Asp Gly Lys Val
 385 390 395 400
 Gly Tyr Met Met Pro Lys Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp
 405 410 415
 Gln Leu Asp Leu Thr Gln Ile Ala Phe Ala Glu Gln Glu Leu Met Leu
 420 425 430
 Lys Asp Lys Lys His Tyr Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu
 435 440 445
 Pro Arg Leu Ala Val Asp Val Ser Ser Leu Pro Met His Ala Gly Asn
 450 455 460
 Ala Thr Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His
 465 470 475 480
 Ile His Val Val Pro Tyr Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr
 485 490 495
 Val Lys Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser
 500 505 510
 Lys Pro Gly His Glu Glu Ser Gly Ser Val Ile Pro Asn Val Thr Pro
 515 520 525
 Leu Asp Lys Arg Ala Gly Met Pro Asn Trp Gln Ile Ile His Ser Ala
 530 535 540
 Glu Glu Val Gln Lys Ala Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp
 545 550 555 560
 Gly Tyr Ile Phe Asp Pro Arg Asp Val Leu Ala Lys Glu Thr Phe Val
 565 570 575
 Trp Lys Asp Gly Ser Phe Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu
 580 585 590
 Arg Thr Ile Asn Lys Ser Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala
 595 600 605
 Gln Glu Leu Leu Ala Lys Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp
 610 615 620
 Lys Pro Lys Glu Lys Gln Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln
 625 630 635 640
 Pro Ser Glu Ala Ser Lys Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp
 645 650 655
 Ser Leu Pro Asp Tyr Gly Leu Asp Arg Ala Thr Leu Glu Asp His Ile
 660 665 670
 Asn Gln Leu Ala Gln Lys Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe
 675 680 685
 Gln Pro Glu Gly Val Gln Phe Tyr Asn Lys Asn Gly Glu Leu Val Thr
 690 695 700
 Tyr Asp Ile Lys Thr Leu Gln Gln Ile Asn Pro
 705 710 715

<210> 17

<211> 77

<212> PRT

<213> Streptococcus

<400> 17

Met His Ser Phe Ser Asn Pro Gly Tyr Pro Tyr Asp Asn Ala Val Thr
 1 5 10 15

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Glu Ala Phe Phe Lys Tyr Leu Lys His Arg Gln Ile Asn Arg Lys His
 20 25 30
 Tyr Gln Asn Ile Lys Gln Val Gln Leu Asp Cys Phe Glu Tyr Ile Glu
 35 40 45
 Asn Phe Tyr Asn Asn Tyr Asn Pro His Thr Ala Asn Leu Gly Leu Thr
 50 55 60
 Pro Asn Gln Lys Glu Glu Asn Tyr Phe Asn Ala Ile Lys
 65 70 75

<210> 18
 <211> 86
 <212> PRT
 <213> Streptococcus

<400> 18
 Met Ala Tyr Tyr Gln Ala Cys Thr Glu Lys Asp Ile Ile Arg Ser Met
 1 5 10 15
 Ser Arg Lys Gly Thr Pro Ala Asp Asn Ala Cys Ile Glu Trp Phe His
 20 25 30
 Thr Val Leu Lys Thr Glu Thr Phe Tyr Phe His Asn Arg Arg Lys Tyr
 35 40 45
 Asn Lys Asp Ser Ile Thr Asn Ile Val Lys Asn Tyr Ile Thr Phe Tyr
 50 55 60
 Asn Glu Thr Arg Ile Gln Gln Arg Leu Asn Asp Gln Ser Pro Val Gln
 65 70 75 80
 Tyr Arg Lys Leu Ile Ala
 85

<210> 19
 <211> 126
 <212> PRT
 <213> Streptococcus

<400> 19
 Met Glu Asn His Phe Ile Tyr Gly Tyr Arg Thr Ile Thr Arg Leu Leu
 1 5 10 15
 Lys Lys Ile His Gly Leu Thr Val Asn Thr Lys Lys Val Tyr Arg Ile
 20 25 30
 Met Lys Asn Asn Gly Trp Leu Cys Arg Thr Arg Thr Lys Lys Val Pro
 35 40 45
 Asn Leu Gly Lys Ala Tyr Tyr Leu Thr Asp Asn Lys Leu Ser Arg Asp
 50 55 60
 Phe His Ala Asp Lys Pro Lys Glu Lys Leu Val Thr Asp Ile Thr Tyr
 65 70 75 80
 Leu Tyr Phe Gly Asn Cys Lys Leu Tyr Leu Ser Ser Ile Met Asn Leu
 85 90 95
 Tyr Asn Arg Glu Ile Ile Ala Tyr Thr Ile Ser Asp Cys Gln Asp Thr
 100 105 110
 Asp Phe Val Leu Asp Thr Leu Asn Gln Leu Lys Leu Pro Lys
 115 120 125

<210> 20
 <211> 96
 <212> PRT
 <213> Streptococcus

36/63

<400> 20

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Met Val Lys Lys Ala Tyr Ser Trp Glu Thr Lys Leu Ala Cys Ile Asp
 1          5          10          15
Met Lys Lys Ala Gly Lys Ser Asn Arg Val Ile Met Glu Thr Leu Gly
          20          25          30
Ile Lys Asn Asn Ser Gln Ile Tyr Thr Trp Met Lys Trp Tyr Glu Asn
          35          40          45
Glu Glu Leu Tyr Arg Phe His Gln Gly Val Gly Lys Gln Tyr Thr Tyr
 50          55          60
Gly Lys Gly Leu Glu His Leu Ser Glu Val Glu Gln Leu Gln Leu Gln
65          70          75          80
Val Asp Leu Leu Lys Lys Tyr Arg Gly Leu Ile Arg Lys Ser Ile Lys
          85          90          95

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<210> 21

<211> 288

<212> PRT

<213> streptococcus

<400> 21

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Ile Arg Tyr Pro Lys Ala Ser Ser Gly Asp Tyr Gly Thr Lys Arg Glu
 1          5          10          15
Ile Ile Thr Ala Asn Lys Asp Lys Tyr Ser Ile Ser Lys Met Cys Arg
          20          25          30
Trp Leu Asn Met Pro His Ser Ser Tyr Tyr Tyr Gln Ala Val Glu Ser
          35          40          45
Val Ser Glu Thr Glu Phe Glu Glu Thr Ile Lys Arg Ile Phe Leu Asp
 50          55          60
Ser Glu Ser Arg Tyr Gly Ser Arg Lys Ile Lys Ile Cys Leu Asn Asn
65          70          75          80
Glu Gly Ile Thr Leu Ser Arg Arg Arg Ile Arg Arg Ile Met Lys Arg
          85          90          95
Leu Asn Leu Val Ser Val Tyr Gln Lys Ala Thr Phe Lys Pro His Ser
          100          105          110
Arg Gly Lys Asn Glu Ala Pro Ile Pro Asn His Leu Asp Arg Gln Phe
          115          120          125
Lys Gln Glu Arg Pro Leu Gln Ala Leu Val Thr Asp Leu Thr Tyr Val
          130          135          140
Arg Val Gly Asn Arg Trp Ala Tyr Val Cys Leu Ile Ile Asp Leu Tyr
145          150          155          160
Asn Arg Glu Ile Ile Gly Leu Ser Leu Gly Trp His Lys Thr Ala Glu
          165          170          175
Leu Val Lys Gln Ala Ile Gln Ser Ile Pro Tyr Ala Leu Thr Lys Val
          180          185          190
Lys Met Phe His Ser Asp Arg Gly Lys Glu Phe Asp Asn Gln Leu Ile
          195          200          205
Asp Glu Ile Leu Glu Ala Phe Gly Ile Thr Arg Ser Leu Ser Gln Ala
210          215          220
Gly Tyr Pro Tyr Asp Asn Ala Val Ala Glu Ser Thr Tyr Arg Ala Phe
225          230          235          240
Lys Ile Glu Phe Val Tyr Gln Glu Thr Phe Gln Leu Leu Glu Glu Leu
          245          250          255
Ala Leu Lys Thr Lys Asp Tyr Val His Trp Trp Asn Tyr His Arg Ile
          260          265          270
His Gly Ser Leu Asn Tyr Gln Thr Pro Met Thr Lys Arg Leu Ile Ala
          275          280          285

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<210> 22
<211> 5058
<212> DNA
<213> streptococcus

<220>
<221> CDS
<222> (1)...(663)

<221> CDS
<222> (763)...(1344)

<221> CDS
<222> (1362)...(1739)

<221> CDS
<222> (2266)...(5058)

<400> 22
aat ttg aaa gca gaa tta tct gta gaa gat gag caa tat aca gca aca      48
Asn Leu Lys Ala Glu Leu Ser Val Glu Asp Glu Gln Tyr Thr Ala Thr
 1              5              10              15

ggt tat ggt aaa tct gct cat ggt tca aca cca caa gaa ggt gtt aat      96
Val Tyr Gly Lys Ser Ala His Gly Ser Thr Pro Gln Glu Gly Val Asn
              20              25              30

ggg gcg act tat tta gct ctt tat cta agt caa ttt gat ttt gaa ggt      144
Gly Ala Thr Tyr Leu Ala Leu Tyr Leu Ser Gln Phe Asp Phe Glu Gly
              35              40              45

cct gct cgt gct ttc tta gat gtt aca gcc aac att att cac gaa gac      192
Pro Ala Arg Ala Phe Leu Asp Val Thr Ala Asn Ile Ile His Glu Asp
              50              55              60

ttc tca ggt gaa aaa ctt gga gta gct tat gaa gat gac tgt atg gga      240
Phe Ser Gly Glu Lys Leu Gly Val Ala Tyr Glu Asp Asp Cys Met Gly
              65              70              75              80

cca ttg agc atg aat gca ggt gtc ttc cag ttt gat gaa act aat gat      288
Pro Leu Ser Met Asn Ala Gly Val Phe Gln Phe Asp Glu Thr Asn Asp
              85              90              95

gat aat act atc gct ctt aat ttc cgt tac cca caa ggg aca gat gct      336
Asp Asn Thr Ile Ala Leu Asn Phe Arg Tyr Pro Gln Gly Thr Asp Ala
              100              105              110

aaa act atc caa act aag ctt gag aaa ctt aac gga gtt gaa aaa gtg      384
Lys Thr Ile Gln Thr Lys Leu Glu Lys Leu Asn Gly Val Glu Lys Val
              115              120              125

act ctt tct gac cat gaa cac aca cca cac tat gta cct atg gac gat      432
Thr Leu Ser Asp His Glu His Thr Pro His Tyr Val Pro Met Asp Asp
              130              135              140

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gaa tta gta tca acc tta cta gct gtc tat gaa aag caa act ggt ctt	480
Glu Leu Val Ser Thr Leu Leu Ala Val Tyr Glu Lys Gln Thr Gly Leu	
145 150 155 160	
aaa gga cat gaa cag gtt att ggt ggt ggg aca ttt ggt cgc tta ctt	528
Lys Gly His Glu Gln Val Ile Gly Gly Gly Thr Phe Gly Arg Leu Leu	
165 170 175	
gaa cgg ggt gtt gca tac ggt gcc atg ttc cca gga gat gaa aac act	576
Glu Arg Gly Val Ala Tyr Gly Ala Met Phe Pro Gly Asp Glu Asn Thr	
180 185 190	
atg cat caa gct aat gag tac atg cct tta gaa aat att ttc cgt tcg	624
Met His Gln Ala Asn Glu Tyr Met Pro Leu Glu Asn Ile Phe Arg Ser	
195 200 205	
gct gct atc tac gca gaa gct atc tat gaa tta atc aaa taaaataatc	673
Ala Ala Ile Tyr Ala Glu Ala Ile Tyr Glu Leu Ile Lys	
210 215 220	
cttaactaa atatgtgac aatgataaag ggtggtgaag acatgaaagt gtctttgcct	733
cttttcataa ggtagattt ggagacttt atg act gac ttg gaa aaa att att	786
Met Thr Asp Leu Glu Lys Ile Ile	
225	
aaa gca ata aaa agt gat tca cag aat caa aat tat aca gaa aat ggt	834
Lys Ala Ile Lys Ser Asp Ser Gln Asn Gln Asn Tyr Thr Glu Asn Gly	
230 235 240 245	
att gat cct ttg ttt gct gct cct aaa aca gct agg atc aat att gtt	882
Ile Asp Pro Leu Phe Ala Ala Pro Lys Thr Ala Arg Ile Asn Ile Val	
250 255 260	
ggc caa gca cct ggt tta aaa act caa gaa gca aga ctc tat tgg aaa	930
Gly Gln Ala Pro Gly Leu Lys Thr Gln Glu Ala Arg Leu Tyr Trp Lys	
265 270 275	
gat aaa tct gga gat cgt cta cgc cag tgg ctt gga gtt gat gaa gag	978
Asp Lys Ser Gly Asp Arg Leu Arg Gln Trp Leu Gly Val Asp Glu Glu	
280 285 290	
aca ttt tac cat tct gga aaa ttt gct gtt tta cct tta gat ttt tat	1026
Thr Phe Tyr His Ser Gly Lys Phe Ala Val Leu Pro Leu Asp Phe Tyr	
295 300 305	
tac cca ggc aaa gga aaa tca gga gat tta ccc cct aga aaa ggt ttt	1074
Tyr Pro Gly Lys Gly Lys Ser Gly Asp Leu Pro Pro Arg Lys Gly Phe	
310 315 320 325	
gcg gag aaa tgg cac cct ctt att tta aaa gaa atg cct aat gtt caa	1122
Ala Glu Lys Trp His Pro Leu Ile Leu Lys Glu Met Pro Asn Val Gln	
330 335 340	
ttg acc ttg cta gtt ggt cag tat gct cag aaa tat tat ctt gga agc	1170
Leu Thr Leu Leu Val Gly Gln Tyr Ala Gln Lys Tyr Tyr Leu Gly Ser	
345 350 355	

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tcc gca cat aaa aat cta aca gaa aca gtt aaa gct tac aaa gac tat	1218
Ser Ala His Lys Asn Leu Thr Glu Thr Val Lys Ala Tyr Lys Asp Tyr	
360 365 370	
cta ccc gat tat tta ccc ctg gtt cac cca tca ccg cga aat caa att	1266
Leu Pro Asp Tyr Leu Pro Leu Val His Pro Ser Pro Arg Asn Gln Ile	
375 380 385	
tgg cta aag aag aat cca tgg ttt gaa aaa gat cta atc gtt gat tta	1314
Trp Leu Lys Lys Asn Pro Trp Phe Glu Lys Asp Leu Ile Val Asp Leu	
390 395 400 405	
caa aag ata gta gca gat att tta aaa gat taaggatagg agttgggt atg	1364
Gln Lys Ile Val Ala Asp Ile Leu Lys Asp Met	
410 415	
aga gat aat cat cta cac acg tat ttt tcc tat gat tgt caa acg gca	1412
Arg Asp Asn His Leu His Thr Tyr Phe Ser Tyr Asp Cys Gln Thr Ala	
420 425 430	
ttt gag gac tat att aat ggt ttt aca ggt gaa ttt atc acg aca gaa	1460
Phe Glu Asp Tyr Ile Asn Gly Phe Thr Gly Glu Phe Ile Thr Thr Glu	
435 440 445	
cat ttt gat tta tca aat cct tac acc ggt caa gac gat gtt cct gat	1508
His Phe Asp Leu Ser Asn Pro Tyr Thr Gly Gln Asp Asp Val Pro Asp	
450 455 460	
tat agt gct tat tgt caa aaa ata gat tat ctt aat cag aaa tat gga	1556
Tyr Ser Ala Tyr Cys Gln Lys Ile Asp Tyr Leu Asn Gln Lys Tyr Gly	
465 470 475 480	
aat cga ttt aaa aaa gga att gaa atc ggt tat ttt aaa gat agg gaa	1604
Asn Arg Phe Lys Lys Gly Ile Glu Ile Gly Tyr Phe Lys Asp Arg Glu	
485 490 495	
tca gat att tta gat tat tta aaa aat aaa gaa ttt gat tta aaa cta	1652
Ser Asp Ile Leu Asp Tyr Leu Lys Asn Lys Glu Phe Asp Leu Lys Leu	
500 505 510	
ttg tca atc cat cat aat ggt agg tat gat tat ctg caa gaa gaa gct	1700
Leu Ser Ile His His Asn Gly Arg Tyr Asp Tyr Leu Gln Glu Glu Ala	
515 520 525	
ctg aaa gta cca aca aag gga gct ttt agc aga tta ctt taatcgtatg	1749
Leu Lys Val Pro Thr Lys Gly Ala Phe Ser Arg Leu Leu	
530 535 540	
gaatttgcca taggccgtgt ggaagcgcac gtttttagctc acttttgatta tggttttcgt	1809
aagttaaact tagatgtaga agatttataaa ccgtttgaaa cgcaattgaa gcgcattttc	1869
ataaagatgt tatctaagggt gttagctttt gaactaaata ccaaattccct ttatctatat	1929
gggaatgaaa aactttatcg ctatgcttta gagataactca aacagcttgg ttgtaaacaa	1989
tactctatag gctctgacgg tcatattcct gaacattttt gttatgaatt tgatagactt	2049
caaggtctgc taaaggacta tcaaattgat gaaaatcatt tgatatgagg aaatttttga	2109
taaaaaagct aggcaatatt gcttagcttt tttgtaatgc tattgatagt tttagtga	2169

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atttcaaaaa aataaagaaa tcatttactt gttgcaagcg cttgcgtaaa ttgttatgat	2229
tttattggta acaattcatt aaaaaaggag aatgat atg aaa aga aaa gac tta	2283
Met Lys Arg Lys Asp Leu	545
ttt ggt gat aaa caa act caa tac acg att aga aag tta agt gtt gga	2331
Phe Gly Asp Lys Gln Thr Gln Tyr Thr Ile Arg Lys Leu Ser Val Gly	550 555 560
gta gct tca gtt aca aca ggg gta tgt att ttt ctt cat agt cca cag	2379
Val Ala Ser Val Thr Thr Gly Val Cys Ile Phe Leu His Ser Pro Gln	565 570 575
gta ttt gct gaa gaa gta agt gtt tct cct gca act aca gcg att gca	2427
Val Phe Ala Glu Glu Val Ser Val Ser Pro Ala Thr Thr Ala Ile Ala	580 585 590 595
gag tcg aat att aat cag gtt gac aac caa caa tct act aat tta aaa	2475
Glu Ser Asn Ile Asn Gln Val Asp Asn Gln Gln Ser Thr Asn Leu Lys	600 605 610
gat gac ata aac tca aac tct gag acg gtt gtg aca ccc tca gat atg	2523
Asp Asp Ile Asn Ser Asn Ser Glu Thr Val Val Thr Pro Ser Asp Met	615 620 625
ccg gat acc aag caa tta gta tca gat gaa act gac act caa aag gga	2571
Pro Asp Thr Lys Gln Leu Val Ser Asp Glu Thr Asp Thr Gln Lys Gly	630 635 640
gtg aca gag ccg gat aag gcg aca agc ctg ctt gaa gaa aat aaa ggt	2619
Val Thr Glu Pro Asp Lys Ala Thr Ser Leu Leu Glu Glu Asn Lys Gly	645 650 655
cct gtt tca gat aaa aat acc tta gat tta aaa gta gca cca tct aca	2667
Pro Val Ser Asp Lys Asn Thr Leu Asp Leu Lys Val Ala Pro Ser Thr	660 665 670 675
ttg caa aat act ccc gac aaa act tct caa gct ata ggt gct cca agc	2715
Leu Gln Asn Thr Pro Asp Lys Thr Ser Gln Ala Ile Gly Ala Pro Ser	680 685 690
cct acc ttg aaa gta gct aat caa gct cca ccg att gaa aat ggt tac	2763
Pro Thr Leu Lys Val Ala Asn Gln Ala Pro Arg Ile Glu Asn Gly Tyr	695 700 705
ttt agg cta cat ctt aaa gaa ttg cct caa ggt cat cct gta gaa agc	2811
Phe Arg Leu His Leu Lys Glu Leu Pro Gln Gly His Pro Val Glu Ser	710 715 720
act gga ctt tgg ata tgg gga gat gtt gat caa ccg tct agt aat tgg	2859
Thr Gly Leu Trp Ile Trp Gly Asp Val Asp Gln Pro Ser Ser Asn Trp	725 730 735
cca aat ggt gct atc cct atg act gat gct aag aaa gat gat tac ggt	2907
Pro Asn Gly Ala Ile Pro Met Thr Asp Ala Lys Lys Asp Asp Tyr Gly	740 745 750 755

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tat tat gtt gat ttt aaa tta tct gaa aaa caa cga aaa caa ata tct	2955
Tyr Tyr Val Asp Phe Lys Leu Ser Glu Lys Gln Arg Lys Gln Ile Ser	
760 765 770	
ttt tta att aat aac aaa gca ggg aca aat tta agc ggc gat cat cat	3003
Phe Leu Ile Asn Asn Lys Ala Gly Thr Asn Leu Ser Gly Asp His His	
775 780 785	
att cca tta tta cga cct gag atg aac caa gtt tgg att gat gaa aag	3051
Ile Pro Leu Leu Arg Pro Glu Met Asn Gln Val Trp Ile Asp Glu Lys	
790 795 800	
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Tyr Gly Ile His Thr Tyr Gln Pro Leu Lys Glu Gly Tyr Val Arg Ile	
805 810 815	
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Asn Tyr Leu Ser Ser Ser Ser Asn Tyr Asp His Leu Ser Ala Trp Leu	
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Phe Lys Asp Val Ala Thr Xaa Ser Thr Thr Trp Pro Asp Gly Ser Asn	
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Phe Val Asn Gln Gly Leu Tyr Gly Arg Tyr Ile Asp Val Ser Leu Lys	
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act aac gcc aaa gag att ggt ttt cta atc tta gat gaa agt aag aca	3291
Thr Asn Ala Lys Glu Ile Gly Phe Leu Ile Leu Asp Glu Ser Lys Thr	
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gga gat gca gtg aaa gtt caa ccc aac gac tat gtt ttt aga gat tta	3339
Gly Asp Ala Val Lys Val Gln Pro Asn Asp Tyr Val Phe Arg Asp Leu	
885 890 895	
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Ala Asn His Asn Gln Ile Phe Val Lys Asp Lys Asp Pro Lys Val Tyr	
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aat aat cct tat tac att gat caa gtg cag cta aag gat gcc caa caa	3435
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920 925 930	
att gat tta aca agt att caa gca agt ttt aca act cta gat ggg gta	3483
Ile Asp Leu Thr Ser Ile Gln Ala Ser Phe Thr Thr Leu Asp Gly Val	
935 940 945	
gat aaa act gaa att tta aaa gaa ttg aaa gtg act gat aaa aat caa	3531
Asp Lys Thr Glu Ile Leu Lys Glu Leu Lys Val Thr Asp Lys Asn Gln	
950 955 960	
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Asn Ala Ile Gln Ile Ser Asp Ile Thr Leu Asp Thr Ser Lys Ser Leu	
965 970 975	

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tct tat aat ggt aac aat gtc atg aca agg caa tct tgg gaa ttt aaa Ser Tyr Asn Gly Asn Asn Val Met Thr Arg Gln Ser Trp Glu Phe Lys 1000 1005 1010	3675
gac caa ctt tat gct tat agt gga aat tta ggt gca gtt ctc aat caa Asp Gln Leu Tyr Ala Tyr Ser Gly Asn Leu Gly Ala Val Leu Asn Gln 1015 1020 1025	3723
gat ggt tca aaa gtt gaa gcc agc ctc tgg tca ccg agt gct gat agt Asp Gly Ser Lys Val Glu Ala Ser Leu Trp Ser Pro Ala Asp Ser 1030 1035 1040	3771
gtc act atg att att tat gac aaa gat aac caa aac agg gtt gta gcg Val Thr Met Ile Ile Tyr Asp Lys Asp Asn Gln Asn Arg Val Val Ala 1045 1050 1055	3819
act acc ccc ctt gtg aaa aat aat aaa ggt gtt tgg cag acg ata ctt Thr Thr Pro Leu Val Lys Asn Asn Lys Gly Val Trp Gln Thr Ile Leu 1060 1065 1070 1075	3867
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 Phe Ser Gly Glu Lys Leu Gly Val Ala Tyr Glu Asp Asp Cys Met Gly
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 Pro Leu Ser Met Asn Ala Gly Val Phe Gln Phe Asp Glu Thr Asn Asp
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 115 120 125
 Thr Leu Ser Asp His Glu His Thr Pro His Tyr Val Pro Met Asp Asp
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 Glu Leu Val Ser Thr Leu Leu Ala Val Tyr Glu Lys Gln Thr Gly Leu
 145 150 155 160
 Lys Gly His Glu Gln Val Ile Gly Gly Gly Thr Phe Gly Arg Leu Leu
 165 170 175
 Glu Arg Gly Val Ala Tyr Gly Ala Met Phe Pro Gly Asp Glu Asn Thr
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<400> 24

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Lys Thr Ala Arg Ile Asn Ile Val Gly Gln Ala Pro Gly Leu Lys Thr
      35           40           45
Gln Glu Ala Arg Leu Tyr Trp Lys Asp Lys Ser Gly Asp Arg Leu Arg
 50           55           60
Gln Trp Leu Gly Val Asp Glu Glu Thr Phe Tyr His Ser Gly Lys Phe
65           70           75           80
Ala Val Leu Pro Leu Asp Phe Tyr Tyr Pro Gly Lys Gly Lys Ser Gly
      85           90           95
Asp Leu Pro Pro Arg Lys Gly Phe Ala Glu Lys Trp His Pro Leu Ile
      100          105          110
Leu Lys Glu Met Pro Asn Val Gln Leu Thr Leu Leu Val Gly Gln Tyr
      115          120          125
Ala Gln Lys Tyr Tyr Leu Gly Ser Ser Ala His Lys Asn Leu Thr Glu
      130          135          140
Thr Val Lys Ala Tyr Lys Asp Tyr Leu Pro Asp Tyr Leu Pro Leu Val
145          150          155          160
His Pro Ser Pro Arg Asn Gln Ile Trp Leu Lys Lys Asn Pro Trp Phe
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Glu Lys Asp Leu Ile Val Asp Leu Gln Lys Ile Val Ala Asp Ile Leu
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Lys Asp

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Glu His Phe Asp Leu Ser Asn Pro Tyr Thr Gly Gln Asp Asp Val Pro
      35           40           45
Asp Tyr Ser Ala Tyr Cys Gln Lys Ile Asp Tyr Leu Asn Gln Lys Tyr
 50           55           60
Gly Asn Arg Phe Lys Lys Gly Ile Glu Ile Gly Tyr Phe Lys Asp Arg
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Glu Ser Asp Ile Leu Asp Tyr Leu Lys Asn Lys Glu Phe Asp Leu Lys
      85           90           95
Leu Leu Ser Ile His His Asn Gly Arg Tyr Asp Tyr Leu Gln Glu Glu
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Phe	Leu	His	Ser	Pro	Gln	Val	Phe	Ala	Glu	Glu	Val	Ser	Val	Ser	Pro
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Ala	Thr	Thr	Ala	Ile	Ala	Glu	Ser	Asn	Ile	Asn	Gln	Val	Asp	Asn	Gln
	50					55				60					
Gln	Ser	Thr	Asn	Leu	Lys	Asp	Asp	Ile	Asn	Ser	Asn	Ser	Glu	Thr	Val
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Val	Thr	Pro	Ser	Asp	Met	Pro	Asp	Thr	Lys	Gln	Leu	Val	Ser	Asp	Glu
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Thr	Asp	Thr	Gln	Lys	Gly	Val	Thr	Glu	Pro	Asp	Lys	Ala	Thr	Ser	Leu
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Leu	Glu	Glu	Asn	Lys	Gly	Pro	Val	Ser	Asp	Lys	Asn	Thr	Leu	Asp	Leu
		115					120					125			
Lys	Val	Ala	Pro	Ser	Thr	Leu	Gln	Asn	Thr	Pro	Asp	Lys	Thr	Ser	Gln
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Ala	Ile	Gly	Ala	Pro	Ser	Pro	Thr	Leu	Lys	Val	Ala	Asn	Gln	Ala	Pro
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Arg	Ile	Glu	Asn	Gly	Tyr	Phe	Arg	Leu	His	Leu	Lys	Glu	Leu	Pro	Gln
				165					170					175	
Gly	His	Pro	Val	Glu	Ser	Thr	Gly	Leu	Trp	Ile	Trp	Gly	Asp	Val	Asp
			180					185					190		
Gln	Pro	Ser	Ser	Asn	Trp	Pro	Asn	Gly	Ala	Ile	Pro	Met	Thr	Asp	Ala
		195					200					205			
Lys	Lys	Asp	Asp	Tyr	Gly	Tyr	Tyr	Val	Asp	Phe	Lys	Leu	Ser	Glu	Lys
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Gln	Arg	Lys	Gln	Ile	Ser	Phe	Leu	Ile	Asn	Asn	Lys	Ala	Gly	Thr	Asn
225					230					235					240
Leu	Ser	Gly	Asp	His	His	Ile	Pro	Leu	Leu	Arg	Pro	Glu	Met	Asn	Gln
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Val	Trp	Ile	Asp	Glu	Lys	Tyr	Gly	Ile	His	Thr	Tyr	Gln	Pro	Leu	Lys
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Glu	Gly	Tyr	Val	Arg	Ile	Asn	Tyr	Leu	Ser	Ser	Ser	Ser	Asn	Tyr	Asp
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His	Leu	Ser	Ala	Trp	Leu	Phe	Lys	Asp	Val	Ala	Thr	Xaa	Ser	Thr	Thr
	290					295					300				
Trp	Pro	Asp	Gly	Ser	Asn	Phe	Val	Asn	Gln	Gly	Leu	Tyr	Gly	Arg	Tyr
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Ile	Asp	Val	Ser	Leu	Lys	Thr	Asn	Ala	Lys	Glu	Ile	Gly	Phe	Leu	Ile
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		340						345					350		
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Lys	Asp	Pro	Lys	Val	Tyr	Asn	Asn	Pro	Tyr	Tyr	Ile	Asp	Gln	Val	Gln
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Leu	Lys	Asp	Ala	Gln	Gln	Ile	Asp	Leu	Thr	Ser	Ile	Gln	Ala	Ser	Phe
385					390					395					400
Thr	Thr	Leu	Asp	Gly	Val	Asp	Lys	Thr	Glu	Ile	Leu	Lys	Glu	Leu	Lys
				405					410					415	
Val	Thr	Asp	Lys	Asn	Gln	Asn	Ala	Ile	Gln	Ile	Ser	Asp	Ile	Thr	Leu
			420					425					430		
Asp	Thr	Ser	Lys	Ser	Leu	Leu	Ile	Ile	Lys	Gly	Asp	Phe	Asn	Pro	Lys
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 Gln Ser Trp Glu Phe Lys Asp Gln Leu Tyr Ala Tyr Ser Gly Asn Leu
 465 470 475 480
 Gly Ala Val Leu Asn Gln Asp Gly Ser Lys Val Glu Ala Ser Leu Trp
 485 490 495
 Ser Pro Ser Ala Asp Ser Val Thr Met Ile Ile Tyr Asp Lys Asp Asn
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 515 520 525
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 545 550 555 560
 Ile Leu Asp Pro Tyr Ala Lys Ser Leu Ala Glu Trp Asp Ser Asn Thr
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 Val Asn Asp Asp Ile Lys Thr Ala Lys Ala Ala Phe Val Asn Pro Ser
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 Gly Arg Gln Asp Ala Val Ile Tyr Glu Ala His Val Arg Asp Phe Thr
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 Ser Asp Arg Ser Leu Asp Gly Lys Leu Lys Asn Gln Phe Gly Thr Phe
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 His Ile Gln Leu Leu Pro Val Leu Ser Tyr Phe Tyr Val Asn Glu Met
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 Asp Lys Ser Arg Ser Thr Ala Tyr Thr Ser Ser Asp Asn Asn Tyr Asn
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 Trp Gly Tyr Asp Pro Gln Ser Tyr Phe Ala Leu Ser Gly Met Tyr Ser
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 Ile His Asp Ile His Lys Arg Gly Met Gly Val Ile Leu Asp Val Val
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 Tyr Asn His Thr Ala Lys Thr Tyr Leu Phe Glu Asp Ile Glu Pro Asn
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 Tyr Tyr His Phe Met Asn Glu Asp Gly Ser Pro Arg Glu Ser Phe Gly
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 770 775 780
 Asp Ser Ile Lys Tyr Leu Thr Ser Glu Phe Lys Val Asp Gly Phe Arg
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 Phe Asp Met Met Gly Asp His Asp Ala Ala Ile Glu Leu Ala Tyr
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 Lys Glu Ala Lys Ala Ile Asn Pro Asn Met Ile Met Ile Gly Glu Gly
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Lys Ala Gln Pro Gly Asn Phe Glu Ala Asp Ser Pro Gly Asp Val Val
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 Lys Ser Ile
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 Arg Ala Gly Asp Thr Val Arg Val His Ala Lys Val Val Glu Gly Thr
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 Arg Glu Arg Ile Gln Ile Phe Glu Gly Val Val Ile Ser Arg Lys Gly
 35 40 45

caa gga atc tca gaa atg tac aca gta cgt aaa att tct ggt ggt atc 193
 Gln Gly Ile Ser Glu Met Tyr Thr Val Arg Lys Ile Ser Gly Gly Ile
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 Arg Ala Leu Gln
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<211> 111

<212> PRT

<213> streptococcus

<400> 28

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 20             25             30
Arg Glu Arg Ile Gln Ile Phe Glu Gly Val Val Ile Ser Arg Lys Gly
 35             40             45
Gln Gly Ile Ser Glu Met Tyr Thr Val Arg Lys Ile Ser Gly Gly Ile
 50             55             60
Gly Val Glu Arg Thr Phe Pro Ile His Thr Pro Arg Val Asp Lys Ile
 65             70             75             80
Glu Val Val Arg Tyr Gly Lys Val Arg Arg Ala Lys Leu Tyr Tyr Leu
 85             90             95
Arg Ala Leu Gln Gly Lys Ala Ala Arg Ile Lys Glu Ile Arg Arg
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<210> 29

<211> 173

<212> PRT

<213> streptococcus

<400> 29

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Thr Asn Lys Tyr Leu Ser Ile Asn Lys Thr Trp Asp Tyr His Phe Asn
 20             25             30
Gln Arg Tyr Leu Pro Thr Lys Asn Lys Ser Ser Ile Arg Asn Ile Pro
 35             40             45
Ile Asp Asn Asp Thr Leu Phe Phe Leu His Glu Phe Thr Lys Asn Lys

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50	55	60
Asn Asp Arg Leu Phe Asp Lys Leu Ser Asn Asn Ala Val Asn Lys Thr		
65	70	75
Ile Arg Lys Ile Thr Gly Arg Glu Val Arg Val His Ser Leu Arg His		
	85	90
Thr Phe Ala Ser Tyr Leu Ile Ser Ile Ser Gln Val Leu Asp His Glu		
	100	105
Asn Leu Asn Ile Thr Leu Glu Val Tyr Ala His Gln Leu Gln Glu Gln		
	115	120
Lys Asp Arg Asn Asp Lys Leu Asn Gln Arg Asn Leu Gly Gln Asn Ser		
	130	135
Ser Lys Pro Leu Phe Thr Cys Asn Glu Tyr Val Pro Cys Arg Asn Arg		
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Thr Ser Asn Tyr Ser Leu Gly Gly Ser Cys Tyr Ile His		
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<211> 389

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<213> streptococcus

<400> 30

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	20	25
Gln Phe Lys Asn Ile Glu Lys Ile Lys Glu Val Glu Glu Lys Ile Phe		
	35	40
Gln Tyr Asp Gly Leu Ala Lys Leu Lys Asp Leu Lys Val Val Ser Gly		
	50	55
Glu Gln Ser Ile Asn Arg Glu Asp Leu Ser Asp Glu Phe Lys Asn Val		
65	70	75
Val Ser Leu Glu Ala Thr Ser Asn Thr Lys Arg Asn Leu Leu Phe Ser		
	85	90
Ser Gly Val Phe Ser Phe Lys Glu Gly Lys Asn Ile Glu Glu Asn Asp		
	100	105
Lys Asn Ser Ile Leu Val His Glu Glu Phe Ala Lys Gln Asn Lys Leu		
	115	120
Lys Leu Gly Asp Glu Ile Asp Leu Glu Leu Leu Asp Thr Glu Lys Ser		
	130	135
Gly Lys Ile Lys Ser His Lys Phe Lys Ile Ile Gly Ile Phe Ser Gly		
145	150	155
Lys Lys Gln Glu Thr Tyr Thr Gly Leu Ser Ser Asp Phe Ser Glu Asn		
	165	170
Met Val Phe Val Asp Tyr Ser Thr Ser Gln Glu Ile Leu Asn Lys Ser		
	180	185
Glu Asn Asn Arg Ile Ala Asn Lys Ile Leu Met Tyr Ser Gly Ser Leu		
	195	200
Glu Ser Thr Glu Leu Ala Leu Asn Lys Leu Lys Asp Phe Lys Ile Asp		
	210	215
Lys Ser Lys Tyr Ser Ile Lys Lys Asp Asn Lys Ala Phe Glu Glu Ser		
225	230	235
Leu Glu Ser Val Ser Gly Ile Lys His Ile Ile Lys Ile Met Thr Tyr		
	245	250
Ser Ile Met Leu Gly Gly Ile Val Val Leu Ser Leu Ile Leu Ile Leu		
	260	265
		270

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Trp Leu Arg Glu Arg Ile Tyr Glu Ile Gly Ile Phe Leu Ser Ile Gly
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 Thr Thr Lys Ile Gln Ile Ile Arg Gln Phe Ile Phe Glu Leu Ile Phe
 290 295 300
 Ile Ser Ile Pro Ser Ile Ile Ser Ser Leu Phe Leu Gly Asn Leu Leu
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 Leu Lys Val Ile Val Glu Gly Phe Ile Asn Ser Glu Asn Ser Met Ile
 325 330 335
 Phe Gly Gly Ser Leu Ile Asn Lys Ser Ser Phe Met Leu Asn Ile Thr
 340 345 350
 Thr Leu Ala Glu Ser Tyr Leu Ile Leu Ile Ser Ile Ile Val Leu Ser
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 <213> streptococcus

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 Phe Tyr Ala Ile Val Gly Lys Ser Gly Thr Gly Lys Ser Thr Leu Leu
 35 40 45
 Ser Leu Leu Ala Gly Leu Asp Lys Val Gln Thr Gly Lys Ile Leu Phe
 50 55 60
 Lys Asn Glu Asp Ile Glu Lys Lys Gly Tyr Ser Asn His Arg Lys Asn
 65 70 75 80
 Asn Ile Ser Leu Val Phe Gln Asn Tyr Asn Leu Ile Asp Tyr Leu Ser
 85 90 95
 Pro Ile Glu Asn Ile Arg Leu Val Asn Lys Ser Val Asp Glu Ser Ile
 100 105 110
 Leu Phe Glu Leu Gly Leu Asp Lys Lys Gln Ile Lys Arg Asn Val Met
 115 120 125
 Lys Leu Ser Gly Gly Gln Gln Gln Arg Val Ala Ile Ala Arg Ala Leu
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 <213> Streptococcus

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 <212> PRT
 <213> Streptococcus

<400> 33

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His	Ala	Asp	Asn	His	Tyr	His	Phe	Phe	Asn	Gly	Lys	Ser	Leu	Ala	Thr	35	40	45	
Phe	Asn	Thr	Asn	Gln	Leu	Ile	Arg	Glu	Val	Val	Tyr	Val	Glu	Ile	Ser	50	55	60	
Leu	Asp	Thr	Met	Ser	Ser	Gly	Glu	His	Asp	Leu	Val	Lys	Val	Asn	Ile	65	70	75	80
Ile	Arg	Pro	Thr	Thr	Glu	His	Thr	Ile	Pro	Thr	Met	Met	Thr	Ala	Ser	85	90	95	
Pro	Tyr	His	Gln	Gly	Ile	Asn	Asp	Pro	Ala	Ala	Asp	Gln	Lys	Thr	Tyr	100	105	110	
Gln	Met	Glu	Gly	Ala	Leu	Ala	Val	Lys	Gln	Pro	Lys	His	Ile	Gln	Val	115	120	125	
Asp	Thr	Lys	Pro	Phe	Lys	Glu	Glu	Val	Lys	His	Pro	Ser	Lys	Leu	Pro	130	135	140	
Ile	Ser	Pro	Ala	Thr	Glu	Ser	Phe	Thr	His	Ile	Asp	Ser	Tyr	Ser	Leu	145	150	155	160
Asn	Asp	Tyr	Phe	Leu	Ser	Arg	Gly	Phe	Ala	Asn	Ile	Tyr	Val	Ser	Gly	165	170	175	
Val	Gly	Thr	Ala	Gly	Ser	Thr	Gly	Phe	Met	Thr	Ser	Gly	Asp	Tyr	Gln	180	185	190	
Gln	Ile	Gln	Ser	Phe	Lys	Ala	Val	Ile	Asp	Trp	Leu	Asn	Gly	Lys	Val	195	200	205	
Thr	Ala	Phe	Thr	Ser	His	Lys	Arg	Asp	Lys	Gln	Val	Lys	Ala	Asp	Trp	210	215	220	
Ser	Asn	Gly	Leu	Val	Ala	Thr	Thr	Gly	Lys	Ser	Tyr	Leu	Gly	Thr	Met	225	230	235	240
Ser	Thr	Gly	Leu	Ala	Thr	Thr	Gly	Val	Glu	Gly	Leu	Lys	Val	Ile	Ile	245	250	255	
Ala	Glu	Ala	Ala	Ile	Ser	Thr	Trp	Tyr	Asp	Tyr	Tyr	Arg	Glu	Asn	Gly	260	265	270	
Leu	Val	Cys	Ser	Pro	Gly	Gly	Tyr	Pro	Gly	Glu	Asp	Leu	Asp	Val	Leu	275	280	285	
Thr	Glu	Leu	Thr	Tyr	Ser	Arg	Asn	Leu	Leu	Ala	Gly	Asp	Tyr	Ile	Lys	290	295	300	
Asn	Asn	Asp	Cys	Tyr	Gln	Ala	Leu	Leu	Asn	Glu	Gln	Ser	Lys	Ala	Ile				

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          325          330          335
Leu Thr His Val Asn Asn Val Lys Ser Arg Val Val Tyr Thr His Gly
          340          345          350
Leu Gln Asp Trp Asn Val Lys Pro Arg His Val Tyr Lys Val Phe Asn
          355          360          365
Ala Leu Pro Gln Thr Ile Lys Lys His Leu Phe Leu His Gln Gly Gln
          370          375          380
His Val Tyr Met His Asn Trp Gln Ser Ile Asp Phe Arg Glu Ser Met
385          390          395          400
Asn Ala Leu Leu Ser Gln Glu Leu Leu Gly Ile Asp Asn His Phe Gln
          405          410          415
Leu Glu Glu Val Ile Trp Gln Asp Asn Thr Thr Glu Gln Thr Trp Gln
          420          425          430
Val Leu Asp Ala Phe Gly Gly Asn His Gln Glu Gln Ile Gly Leu Gly
          435          440          445
Asp Ser Lys Lys Leu Ile Asp Asn His Tyr Asp Lys Glu Ala Phe Asp
          450          455          460
Thr Tyr Cys Lys Asp Phe Asn Val Phe Lys Asn Asp Leu Phe Lys Gly
465          470          475          480
Asn Asn Lys Thr Asn Gln Ile Thr Ile Asn Leu Pro Leu Lys Lys Asn
          485          490          495
Tyr Leu Leu Asn Gly Gln Cys Lys Leu His Leu Arg Val Lys Thr Ser
          500          505          510
Asp Lys Lys Ala Ile Leu Ser Ala Gln Ile Leu Asp Tyr Gly Pro Lys
          515          520          525
Lys Arg Phe Lys Asp Thr Pro Thr Ile Lys Phe Leu Asn Ser Leu Asp
          530          535          540
Asn Gly Lys Asn Phe Ala Arg Glu Ala Leu Arg Glu Leu Pro Phe Thr
545          550          555          560
Lys Asp His Tyr Arg Val Ile Ser Lys Gly Val Leu Asn Leu Gln Asn
          565          570          575
Arg Thr Asp Leu Leu Thr Ile Glu Ala Ile Glu Pro Glu Gln Trp Phe
          580          585          590
Asp Ile Glu Phe Ser Leu Gln Pro Ser Ile Tyr Gln Leu Ser Lys Gly
          595          600          605
Asp Asn Leu Arg Ile Ile Leu Tyr Thr Thr Asp Phe Glu His Thr Ile
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<211> 119
<212> PRT
<213> Streptococcus

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<400> 34
Met Lys Leu Leu Thr Lys Glu Arg Phe Asp Asp Ser Gln His Phe Trp
1          5          10          15
Tyr Gln Ile Asn Leu Leu Gln Glu Ser Asn Phe Gly Ala Val Phe Asp
          20          25          30
His Asp Asn Lys Asn Ile Pro Gln Val Val Ala Thr Ile Val Asp Asp
          35          40          45

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Leu Gln Gly Ser Gly Ser Ser Asn His Phe Trp Tyr Phe Gly Asn Thr
 50 55 60
 Thr Asp Thr Ser Ile Leu Met Ile Ala His Leu Asn Arg Lys Phe Tyr
 65 70 75 80
 Ile Gln Val Asn Leu Lys Asp Phe Asp Phe Ala Leu Asn Leu Ile Ala
 85 90 95
 Ile Asn Asn Trp Lys Ser Leu Leu Gln Thr Gln Leu Glu Ala Leu Asn
 100 105 110
 Asp Thr Leu Ala Ile Phe Gln
 115

<210> 35
 <211> 326
 <212> PRT
 <213> Streptococcus

<400> 35
 Met Ser Ser Tyr Trp Asn Asn Tyr Pro Glu Leu Lys Lys Asn Ile Asp
 1 5 10 15
 Glu Thr Asn Gln Leu Ile Gln Glu Arg Ile Gln Val Arg Asn Lys Asp
 20 25 30
 Ile Glu Ala Ala Leu Ser Gln Leu Thr Ala Ala Gly Gly Lys Gln Leu
 35 40 45
 Arg Pro Ala Phe Phe Tyr Leu Phe Ser Gln Leu Gly Asn Lys Glu Asn
 50 55 60
 Gln Asp Thr Gln Gln Leu Lys Lys Ile Ala Ala Ser Leu Glu Ile Leu
 65 70 75 80
 His Val Ala Thr Leu Ile His Asp Asp Val Ile Asp Asp Ser Pro Leu
 85 90 95
 Arg Arg Gly Asn Met Thr Ile Gln Ser Lys Phe Gly Lys Asp Ile Ala
 100 105 110
 Val Tyr Thr Gly Asp Leu Leu Phe Thr Val Phe Phe Asp Leu Ile Leu
 115 120 125
 Glu Ser Met Thr Asp Thr Pro Phe Met Arg Ile Asn Ala Lys Ser Met
 130 135 140
 Arg Lys Ile Leu Met Gly Glu Leu Asp Gln Met His Leu Arg Tyr Asn
 145 150 155 160
 Gln Gln Gln Gly Ile His His Tyr Leu Arg Ala Ile Ser Gly Lys Thr
 165 170 175
 Ala Glu Leu Phe Lys Leu Ala Ser Lys Glu Gly Ala Tyr Phe Gly Gly
 180 185 190
 Ala Glu Lys Glu Val Val Arg Leu Ala Gly His Ile Gly Phe Asn Ile
 195 200 205
 Gly Met Thr Phe Gln Ile Leu Asp Asp Ile Leu Asp Tyr Thr Ala Asp
 210 215 220
 Lys Lys Thr Phe Asn Lys Pro Val Leu Glu Asp Leu Thr Gln Gly Val
 225 230 235 240
 Tyr Ser Leu Pro Leu Leu Leu Ala Ile Glu Glu Asn Pro Asp Ile Phe
 245 250 255
 Lys Pro Ile Leu Asp Lys Lys Thr Asp Met Ala Thr Glu Asp Met Glu
 260 265 270
 Lys Ile Ala Tyr Leu Val Val Ser His Arg Gly Val Asp Lys Ala Arg
 275 280 285
 His Leu Ala Arg Lys Phe Thr Glu Lys Ala Ile Ser Asp Ile Asn Lys
 290 295 300
 Leu Pro Gln Asn Ser Ala Lys Lys Gln Leu Leu Gln Leu Thr Asn Tyr

305	310	315	320
Leu	Leu	Lys	Arg
	Lys	Ile	
	325		

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<210> 36
<211> 247
<212> PRT
<213> Streptococcus
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<400> 36															
Leu 1	Pro	Asn	Lys	Pro 5	Tyr	Asp	Phe	Ser	Val 10	Lys	Asn	Leu	Ser	Phe 15	Gln
Tyr	Lys	Pro	Gln 20	Glu	Lys	Trp	Val	Leu 25	His	His	Leu	Asp	Leu 30	Asp	Ile
Lys	Glu	Gly 35	Glu	Lys	Ile	Ala	Ile 40	Leu	Gly	Arg	Ser	Gly 45	Ser	Gly	Lys
Ser	Thr 50	Leu	Ala	Ser	Leu	Leu 55	Arg	Gly	Asp	Leu	Lys 60	Ala	Ser	Gln	Gly
Lys 65	Ile	Thr	Leu	Gly	Gly 70	Ala	Asp	Val	Ser	Ile 75	Val	Gly	Asp	Cys 80	Ile
Ser	Asn	Tyr	Ile	Gly 85	Val	Ile	Gln	Gln	Ala 90	Pro	Tyr	Leu	Phe 95	Asn	Thr
Thr	Leu	Leu	Asn 100	Asn	Ile	Arg	Ile	Gly 105	Asn	Gln	Asp	Ala	Ser 110	Glu	Glu
Asp	Val	Trp 115	Lys	Val	Leu	Glu	Arg 120	Val	Gly	Leu	Lys	Glu 125	Met	Val	Thr
Asp	Leu	Ser 130	Asp	Gly	Leu	Tyr	Thr 135	Met	Val	Asp	Glu 140	Ala	Gly	Leu	Arg
Phe 145	Ser	Gly	Gly	Glu	Arg 150	His	Arg	Ile	Ala	Leu 155	Ala	Arg	Ile	Leu 160	Leu
Lys	Asp	Val	Pro	Ile 165	Val	Ile	Leu	Asp	Glu 170	Pro	Thr	Val	Gly	Leu 175	Asp
Pro	Ile	Thr 180	Glu	Gln	Ala	Leu	Leu	Arg 185	Val	Phe	Met	Lys	Glu 190	Leu	Glu
Gly	Lys	Thr 195	Leu	Val	Trp	Ile	Thr 200	His	His	Leu	Lys	Gly 205	Ile	Glu	His
Ala	Asp	Arg 210	Ile	Leu	Phe	Ile	Glu 215	Asn	Gly	Gln	Leu	Glu 220	Leu	Glu	Gly
Ser 225	Pro	Gln	Glu	Leu	Ser 230	Gln	Ser	Ser	Gln	Arg 235	Tyr	Arg	Gln	Leu	Lys 240
Ala	Ala	Asp	Asp	Gly 245	Asp	Leu									

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<210> 37
<211> 3480
<212> DNA
<213> Streptococcus
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<400> 37						
aattctattt	ggaggttttt	cttgaataaa	tggtagtta	aggcaagttc	cttagttggt	60
ttaggttgta	tggttttatc	tgcgggttcc	cgagttttag	cggatactta	tgtccgtcca	120
attgataatg	gtagaattac	aacaggtttc	aatggttatc	ctggacattg	tggggtggat	180
tatgctgttc	cgactggaac	gattattagg	gcagtggcag	atgggtactgt	gaaatttgca	240
ggagctggag	ccaacttttc	tggatgaca	gacttagcac	gaaatttgtgt	catgattcaa	300
catgcgggat	gaatgcatag	tgtttacgct	catatgtcac	gtgtggtggc	taggactggg	360
qaaaaagtca	aacaaggaga	tatcatcggt	tacgtaggag	caactggtat	ggcgacggga	420

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cctcaccttc attttgaatt tttaccagct aaccctaatt ttcaaaatgg tttccatgga 480
cgtatcaatc caacgtcact aattgctaac gttgcgacct ttagtggaag aacgcaagca 540
tcagctccaa gcattaagcc attacaatca gctcctgtac agaataatc tagtaaatga 600
aaagtgtatc gagtagatga attacaaaag gttaaatggtg tttgggttagt caaaaaatac 660
accctaacgc cgactgggtt tgattggaac gataatggta taccagcatc agaaattgat 720
gagggttgatg ctaatggtaa tttgacagct gaccagggttc ttcaaaaagg tgggtacttt 780
atcctttaatc ctaaaactct taagactgta gaaaaaccca tccaagggaac agctgggtta 840
acttgggcta agacacgctt tgctaattgt agttcagttt ggcttcgcgt tgacaacagt 900
caagaactgc tttacaaata gtttgaggta ttgattcatt gttttaaatg acagttttgt 960
tactaactaa gtacaatttc tttaaaccgt ctgaaaataa ttttatagtc cagtaaagtg 1020
tgatattata gtctcggact aataaaaagg aaataggaat tgaagcaatg aaaaatgaata 1080
aaaaggtact attgacatcg acaatggcag ctctcgctatt atcagtcgca agtgttcaag 1140
cacaagaaac agatacgacg tggacagcac gtactgtttc agaggtaaag gctgatttgg 1200
taaagcaaga caataaatca tcatatactg tgaaatatgg tgatacacta agcgttattt 1260
cagaagcaat gtcaattgat atgaatgtct tagcaaaaat taataacatt gcagatatca 1320
atccttattta tcctgagaca acactgacag taacttacga tcagaagagt catactgcca 1380
cttcaatgaa aatagaaaca ccagcaacaa atgctgctgg tcaaacacaa gctactgtgg 1440
atltgaaaac caatcaagtt tctgttgacg accaaaaagt ttctctcaat acaatttcgg 1500
aagggtatgac accagaagca gcaacaacga ttgtttcgcc aatgaagaca tattcttctg 1560
cgccagcttt gaaatcaaaa gaagtattag cacaagagca agctgttagt caagcagcag 1620
ctaatagaac ggtatcaaca gctcctgtga agtcgattac ttcagaagtt ccagcagcta 1680
aagaggaagt taaccact cagacgtcag tcagtcagtc aacaacagta tcaccagctt 1740
ctgttgccgc tgaaacacca gctccagtag ctaaagtagc accggtaaga actgtagcag 1800
cccctagagt ggcaagtgtt aaagtagtca ctctaaagt agaaactggg gcatcaccag 1860
agcatgtatc agctccagca gttcctgtga ctacgacttc aacagctaca gacagtaagt 1920
tacaagcgac tgaagttaag agcgttccgg tagcacaaaa agctccaaca gcaacaccgg 1980
tagcacaacc agcttcaaca acaaatgcag tagctgcaca tcctgaaaat gcagggtccc 2040
aacctcatgt tgcagcttat aaagaaaaag tagcgtcaac ttatggagt aatgaattca 2100
gtacataccg tgcaggtgat ccaggtgatc atggtaaagg tttagcagtc gactttattg 2160
taggtaaaaa ccaagcactt ggtaataag ttgcacagta ctctacacaa aatatggcag 2220
caataacat ttcatatgtt atctggcaac aaaagtttta ctcaaataca aatagtattt 2280
atggacctgc taatacttgg aatgcaatgc cagatcgtgg tggcgttact gccaacatt 2340
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ttaagctata agtctgaac tactttcacg ttaaccgtga ctaaatcaaa acgttaaaac 2520
taaaatctaa gtctgtaaag attattgaaa acgctttaaa aacagatata ataaggtttg 2580
tagatatcta aaattaaaaa agataaggaa gtgagaatat gccacatcta agtaaaagag 2640
cttttaaaaa gcaataaaaa aatggcatta ttgtgtcatg tcaagctttg cctggggagc 2700
ctcttttatac tgaagtggga ggtgttatgc ctcttttagc tttggcagct caagaagcag 2760
gagcgggttg tataagagcc aatagtgtcc gcgacattaa ggaaattcaa gaagtacta 2820
atltacctat catcggcatt attaaacgtg aatatcctcc acaagaacca tttatcactg 2880
ctacgatgac agaggtggat caattagcta gtttagatat tgcagtaata gccttagatt 2940
gtacacttag agagcgtcat gatggtttga gtgtagctga gtttattcaa aagataaaaag 3000
ggaaatatcc tgaacagttg ctaatggctg atataagtac ttttgaagaa ggtaaaaatg 3060
cttttgaagc aggagttgat tttgtgggta caactctatc tggatacaca gattacagcc 3120
gccaagaaga aggaccgat atagaactcc ttaataagct ttgtcaagcc ggtatagatg 3180
tgattgcgga aggtaaaatt catactccta agcaagctaa tgaaattaat catataggtg 3240
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tctcaggact tagttaaaag tgttactcaa aaatcaaaat caaaataaaa aagggggaata 3360
gttatgagta tcaaaaaaag tgtgattgggt ttttgcctcg gagctgcagc attatcaatg 3420
tttgcttgtg tagacagtag tcaatctgtt atggctgccg agaaggataa agtcgaaatt 3480

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<210> 38

<211> 306

<212> PRT

<213> Streptococcus

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<400> 38

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Asn Ser Ile Trp Arg Phe Phe Leu Asn Lys Trp Leu Val Lys Ala Ser
 1           5           10           15
Ser Leu Val Val Leu Gly Gly Met Val Leu Ser Ala Gly Ser Arg Val
 20           25           30
Leu Ala Asp Thr Tyr Val Arg Pro Ile Asp Asn Gly Arg Ile Thr Thr
 35           40           45
Gly Phe Asn Gly Tyr Pro Gly His Cys Gly Val Asp Tyr Ala Val Pro
 50           55           60
Thr Gly Thr Ile Ile Arg Ala Val Ala Asp Gly Thr Val Lys Phe Ala
 65           70           75           80
Gly Ala Gly Ala Asn Phe Ser Trp Met Thr Asp Leu Ala Gly Asn Cys
 85           90           95
Val Met Ile Gln His Ala Asp Gly Met His Ser Gly Tyr Ala His Met
 100          105          110
Ser Arg Val Val Ala Arg Thr Gly Glu Lys Val Lys Gln Gly Asp Ile
 115          120          125
Ile Gly Tyr Val Gly Ala Thr Gly Met Ala Thr Gly Pro His Leu His
 130          135          140
Phe Glu Phe Leu Pro Ala Asn Pro Asn Phe Gln Asn Gly Phe His Gly
 145          150          155          160
Arg Ile Asn Pro Thr Ser Leu Ile Ala Asn Val Ala Thr Phe Ser Gly
 165          170          175
Lys Thr Gln Ala Ser Ala Pro Ser Ile Lys Pro Leu Gln Ser Ala Pro
 180          185          190
Val Gln Asn Gln Ser Ser Lys Leu Lys Val Tyr Arg Val Asp Glu Leu
 195          200          205
Gln Lys Val Asn Gly Val Trp Leu Val Lys Asn Asn Thr Leu Thr Pro
 210          215          220
Thr Gly Phe Asp Trp Asn Asp Asn Gly Ile Pro Ala Ser Glu Ile Asp
 225          230          235          240
Glu Val Asp Ala Asn Gly Asn Leu Thr Ala Asp Gln Val Leu Gln Lys
 245          250          255
Gly Gly Tyr Phe Ile Phe Asn Pro Lys Thr Leu Lys Thr Val Glu Lys
 260          265          270
Pro Ile Gln Gly Thr Ala Gly Leu Thr Trp Ala Lys Thr Arg Phe Ala
 275          280          285
Asn Gly Ser Ser Val Trp Leu Arg Val Asp Asn Ser Gln Glu Leu Leu
 290          295          300
Tyr Lys
305

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<210> 39

<211> 434

<212> PRT

<213> Streptococcus

<400> 39

```

Met Lys Met Asn Lys Lys Val Leu Leu Thr Ser Thr Met Ala Ala Ser
 1           5           10           15
Leu Leu Ser Val Ala Ser Val Gln Ala Gln Glu Thr Asp Thr Thr Trp
 20           25           30
Thr Ala Arg Thr Val Ser Glu Val Lys Ala Asp Leu Val Lys Gln Asp
 35           40           45
Asn Lys Ser Ser Tyr Thr Val Lys Tyr Gly Asp Thr Leu Ser Val Ile

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50	55	60
Ser Glu Ala Met Ser Ile Asp Met Asn Val Leu Ala Lys Ile Asn Asn		
65	70	75
Ile Ala Asp Ile Asn Leu Ile Tyr Pro Glu Thr Thr Leu Thr Val Thr		80
	85	90
Tyr Asp Gln Lys Ser His Thr Ala Thr Ser Met Lys Ile Glu Thr Pro		95
	100	105
Ala Thr Asn Ala Ala Gly Gln Thr Thr Ala Thr Val Asp Leu Lys Thr		110
	115	120
Asn Gln Val Ser Val Ala Asp Gln Lys Val Ser Leu Asn Thr Ile Ser		125
	130	135
Glu Gly Met Thr Pro Glu Ala Ala Thr Thr Ile Val Ser Pro Met Lys		140
145	150	155
Thr Tyr Ser Ser Ala Pro Ala Leu Lys Ser Lys Glu Val Leu Ala Gln		160
	165	170
Glu Gln Ala Val Ser Gln Ala Ala Ala Asn Glu Gln Val Ser Thr Ala		175
	180	185
Pro Val Lys Ser Ile Thr Ser Glu Val Pro Ala Ala Lys Glu Glu Val		190
	195	200
Lys Pro Thr Gln Thr Ser Val Ser Gln Ser Thr Thr Val Ser Pro Ala		205
	210	215
Ser Val Ala Ala Glu Thr Pro Ala Pro Val Ala Lys Val Ala Pro Val		220
225	230	235
Arg Thr Val Ala Ala Pro Arg Val Ala Ser Val Lys Val Val Thr Pro		240
	245	250
Lys Val Glu Thr Gly Ala Ser Pro Glu His Val Ser Ala Pro Ala Val		255
	260	265
Pro Val Thr Thr Thr Ser Thr Ala Thr Asp Ser Lys Leu Gln Ala Thr		270
	275	280
Glu Val Lys Ser Val Pro Val Ala Gln Lys Ala Pro Thr Ala Thr Pro		285
	290	295
Val Ala Gln Pro Ala Ser Thr Thr Asn Ala Val Ala Ala His Pro Glu		300
305	310	315
Asn Ala Gly Leu Gln Pro His Val Ala Ala Tyr Lys Glu Lys Val Ala		320
	325	330
Ser Thr Tyr Gly Val Asn Glu Phe Ser Thr Tyr Arg Ala Gly Asp Pro		335
	340	345
Gly Asp His Gly Lys Gly Leu Ala Val Asp Phe Ile Val Gly Lys Asn		350
	355	360
Gln Ala Leu Gly Asn Glu Val Ala Gln Tyr Ser Thr Gln Asn Met Ala		365
	370	375
Ala Asn Asn Ile Ser Tyr Val Ile Trp Gln Gln Lys Phe Tyr Ser Asn		380
385	390	395
Thr Asn Ser Ile Tyr Gly Pro Ala Asn Thr Trp Asn Ala Met Pro Asp		400
	405	410
Arg Gly Gly Val Thr Ala Asn His Tyr Asp His Val His Val Ser Phe		415
	420	425
Asn Lys		430

<210> 40

<211> 232

<212> PRT

<213> Streptococcus

<400> 40

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Met Pro His Leu Ser Lys Glu Ala Phe Lys Lys Gln Ile Lys Asn Gly
 1 5 10 15
 Ile Ile Val Ser Cys Gln Ala Leu Pro Gly Glu Pro Leu Tyr Thr Glu
 20 25 30
 Ser Gly Gly Val Met Pro Leu Leu Ala Leu Ala Ala Gln Glu Ala Gly
 35 40 45
 Ala Val Gly Ile Arg Ala Asn Ser Val Arg Asp Ile Lys Glu Ile Gln
 50 55 60
 Glu Val Thr Asn Leu Pro Ile Ile Gly Ile Ile Lys Arg Glu Tyr Pro
 65 70 75 80
 Pro Gln Glu Pro Phe Ile Thr Ala Thr Met Thr Glu Val Asp Gln Leu
 85 90 95
 Ala Ser Leu Asp Ile Ala Val Ile Ala Leu Asp Cys Thr Leu Arg Glu
 100 105 110
 Arg His Asp Gly Leu Ser Val Ala Glu Phe Ile Gln Lys Ile Lys Gly
 115 120 125
 Lys Tyr Pro Glu Gln Leu Leu Met Ala Asp Ile Ser Thr Phe Glu Glu
 130 135 140
 Gly Lys Asn Ala Phe Glu Ala Gly Val Asp Phe Val Gly Thr Thr Leu
 145 150 155 160
 Ser Gly Tyr Thr Asp Tyr Xaa Arg Gln Glu Gly Pro Asp Ile Glu
 165 170 175
 Leu Leu Asn Lys Leu Cys Gln Ala Gly Ile Asp Val Ile Ala Glu Gly
 180 185 190
 Lys Ile His Thr Pro Lys Gln Ala Asn Glu Ile Asn His Ile Gly Val
 195 200 205
 Ala Gly Ile Val Val Gly Gly Ala Ile Thr Arg Pro Lys Glu Ile Ala
 210 215 220
 Glu Arg Phe Ile Ser Gly Leu Ser
 225 230

<210> 41

<211> 39

<212> PRT

<213> Streptococcus

<400> 41

Met Ser Ile Lys Lys Ser Val Ile Gly Phe Cys Leu Gly Ala Ala Ala
 1 5 10 15
 Leu Ser Met Phe Ala Cys Val Asp Ser Ser Gln Ser Val Met Ala Ala
 20 25 30
 Glu Lys Asp Lys Val Glu Ile
 35

<210> 42

<211> 1305

<212> DNA

<213> Streptococcus

<400> 42

atgaaaatga ataaaaagggt actattgaca tcgacaatgg cagcttcgct attatcagtc 60
 gcaagtgttc aagcacaaga aacagatacg acgtggacag cacgtactgt ttcagaggta 120
 aaggctgatt tggtaaagca agacaataaa tcatcatata ctgtgaaata tggatgataca 180
 ctaagcgtta tttcagaagc aatgtcaatt gatatgaatg tcttagcaaa aattaataac 240
 attgcagata tcaatcttat ttatcctgag acaacactga cagtaactta cgatcagaag 300
 agtcatactg ccacttcaat gaaaaatagaa acaccagcaa caaatgctgc tggcacaaca 360

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acagctactg tggatttgaa aaccaatcaa gtttctgttg cagacaaaa agtttctctc 420
aatacaattt cggaaggatg gacaccagaa gcagcaacaa cgattgtttc gccaatgaag 480
acataattctt ctgcgccagc ttgaaatca aaagaagtat tagcacaaga gcaagctggt 540
agtcaagcag cagctaatag acaggatca acagctcctg tgaagtcgat tacttcagaa 600
gttccagcag ctaaagagga agttaacca actcagacgt cagtcagtca gtcaacaaca 660
gtatcaccag cttctgttgc cgctgaaaca ccagctccag tagctaaagt agcaccggta 720
agaactgtag cagcccttag agtggcaagt gttaaagttag tcactcctaa agtagaaact 780
gggtgcatcac cagagcatgt atcagctcca gcagttcctg tgactacgac ttcaacagct 840
acagacagta agttacaagc gactgaagtt aagagcggtc cggtagcaca aaaagctcca 900
acagcaacac cggtagcaca accagcttca acaacaaatg cagtagctgc acatcctgaa 960
aatgcagggc tccaacctca tgttcagct tataaagaaa aagtagcgct aacttatgga 1020
gttaatgaat tcagtacata ccgtgcaggt gatccaggtg atcatggtaa aggttttagca 1080
gtcgacttta ttgtaggtaa aaaccaagca cttggtaatg aagttgcaca gtactctaca 1140
caaaatatgg cagcaataa catttcatat gttatctggc aacaaaagtt ttactcaaat 1200
acaaatagta tttatggacc tgctaatact tggaaatgca tgccagatcg tggtagcggt 1260
actgccacc attatgacca tgttcacgta tcatttaaca aataa 1305

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<210> 43

<211> 1230

<212> DNA

<213> Streptococcus

<400> 43

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caagaaacag atacgacgtg gacagcacgt actgtttcag aggtaaaggc tgatttggtg 60
aagcaagaca ataaatcatc atatactgtg aaatatggtg atacactaag cgttatttca 120
gaagcaatgt caattgatat gaatgtctta gcaaaaatta ataacattgc agatatcaat 180
cttattttatc ctgagacaac actgacagta acttacgatc agaagagtca tactgccact 240
tcaatgaaaa tagaaacacc agcaacaaat gctgctggtc aaacaacagc tactgtggat 300
ttgaaaacca atcaagtttc tgttcagac caaaaagttt ctctcaatac aatttcggaa 360
gggatgacac cagaagcagc aacaacgatt gtttcgcaa tgaagacata ttcttctgcg 420
ccagctttga aatcaaaaga agtattagca caagagcaag ctgttagtca agcagcagct 480
aatgaacagg tatcaacagc tcctgtgaag tcgattactt cagaagttcc agcagctaaa 540
gaggaagtta aaccaactca gacgtcagtc agtcagtcaa caacagtatc accagcttct 600
gttgccgctg aaacaccagc tccagtagct aaagtagcac cggtaaagaa tgtagcagcc 660
cctagagtgg caagtgttaa agtagtcact cctaaagtag aaactgggtg atcaccagag 720
catgtatcag ctccagcagt tcctgtgact acgacttcaa cagctacaga cagtaagtta 780
caagcgactg aagttaagag cgttcgggta gcacaaaaag ctccaacagc aacaccggta 840
gcacaaccag cttcaacaac aaatgcagta gctgcacatc ctgaaaatgc agggctccaa 900
cctcatgttg cagcttataa agaaaaagta gcgtcaactt atggagttaa tgaattcagt 960
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ggtaaaaacc aagcacttgg taatgaagtt gcacagtact ctacacaaaa tatggcagca 1080
aataacattt catatgttat ctggcaacaa aagttttact caaatacaaa tagtatttat 1140
ggacctgcta atacttggaa tgcaatgcc aatcggtg gcgttactgc caaccattat 1200
gaccatgttc acgtatcatt taacaaataa 1230

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<210> 44

<211> 409

<212> PRT

<213> Streptococcus

<400> 44

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Gln Glu Thr Asp Thr Thr Trp Thr Ala Arg Thr Val Ser Glu Val Lys
 1             5             10             15
Ala Asp Leu Val Lys Gln Asp Asn Lys Ser Ser Tyr Thr Val Lys Tyr
      20             25             30
Gly Asp Thr Leu Ser Val Ile Ser Glu Ala Met Ser Ile Asp Met Asn

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      35              40              45
Val Leu Ala Lys Ile Asn Asn Ile Ala Asp Ile Asn Leu Ile Tyr Pro
  50              55              60
Glu Thr Thr Leu Thr Val Thr Tyr Asp Gln Lys Ser His Thr Ala Thr
  65              70              75              80
Ser Met Lys Ile Glu Thr Pro Ala Thr Asn Ala Ala Gly Gln Thr Thr
      85              90              95
Ala Thr Val Asp Leu Lys Thr Asn Gln Val Ser Val Ala Asp Gln Lys
  100              105              110
Val Ser Leu Asn Thr Ile Ser Glu Gly Met Thr Pro Glu Ala Ala Thr
  115              120              125
Thr Ile Val Ser Pro Met Lys Thr Tyr Ser Ser Ala Pro Ala Leu Lys
  130              135              140
Ser Lys Glu Val Leu Ala Gln Glu Gln Ala Val Ser Gln Ala Ala Ala
  145              150              155              160
Asn Glu Gln Val Ser Thr Ala Pro Val Lys Ser Ile Thr Ser Glu Val
      165              170              175
Pro Ala Ala Lys Glu Glu Val Lys Pro Thr Gln Thr Ser Val Ser Gln
      180              185              190
Ser Thr Thr Val Ser Pro Ala Ser Val Ala Ala Glu Thr Pro Ala Pro
  195              200              205
Val Ala Lys Val Ala Pro Val Arg Thr Val Ala Ala Pro Arg Val Ala
  210              215              220
Ser Val Lys Val Val Thr Pro Lys Val Glu Thr Gly Ala Ser Pro Glu
  225              230              235              240
His Val Ser Ala Pro Ala Val Pro Val Thr Thr Thr Ser Thr Ala Thr
      245              250              255
Asp Ser Lys Leu Gln Ala Thr Glu Val Lys Ser Val Pro Val Ala Gln
      260              265              270
Lys Ala Pro Thr Ala Thr Pro Val Ala Gln Pro Ala Ser Thr Thr Asn
      275              280              285
Ala Val Ala Ala His Pro Glu Asn Ala Gly Leu Gln Pro His Val Ala
  290              295              300
Ala Tyr Lys Glu Lys Val Ala Ser Thr Tyr Gly Val Asn Glu Phe Ser
  305              310              315              320
Thr Tyr Arg Ala Gly Asp Pro Gly Asp His Gly Lys Gly Leu Ala Val
      325              330              335
Asp Phe Ile Val Gly Lys Asn Gln Ala Leu Gly Asn Glu Val Ala Gln
      340              345              350
Tyr Ser Thr Gln Asn Met Ala Ala Asn Asn Ile Ser Tyr Val Ile Trp
      355              360              365
Gln Gln Lys Phe Tyr Ser Asn Thr Asn Ser Ile Tyr Gly Pro Ala Asn
  370              375              380
Thr Trp Asn Ala Met Pro Asp Arg Gly Gly Val Thr Ala Asn His Tyr
  385              390              395              400
Asp His Val His Val Ser Phe Asn Lys
      405

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